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(54) Title: TREATMENT FOR ATHEROSCLEROSIS AND OTHER CARDIOVASCULAR AND INFLAMMATORY DISEASES

(57) Abstract

Dithiocarboxylates, including dithiocarbamates, block the induced expression of the endothelial cell surface adhesion molecule VCAM-1, and are therefore useful in the treatment of cardiovascular disease, including atherosclerosis, as well as noncardiovascular inflammatory diseases that are mediated by VCAM-1.

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Treatment for Atherosclerosis  
and Other Cardiovascular and Inflammatory Diseases

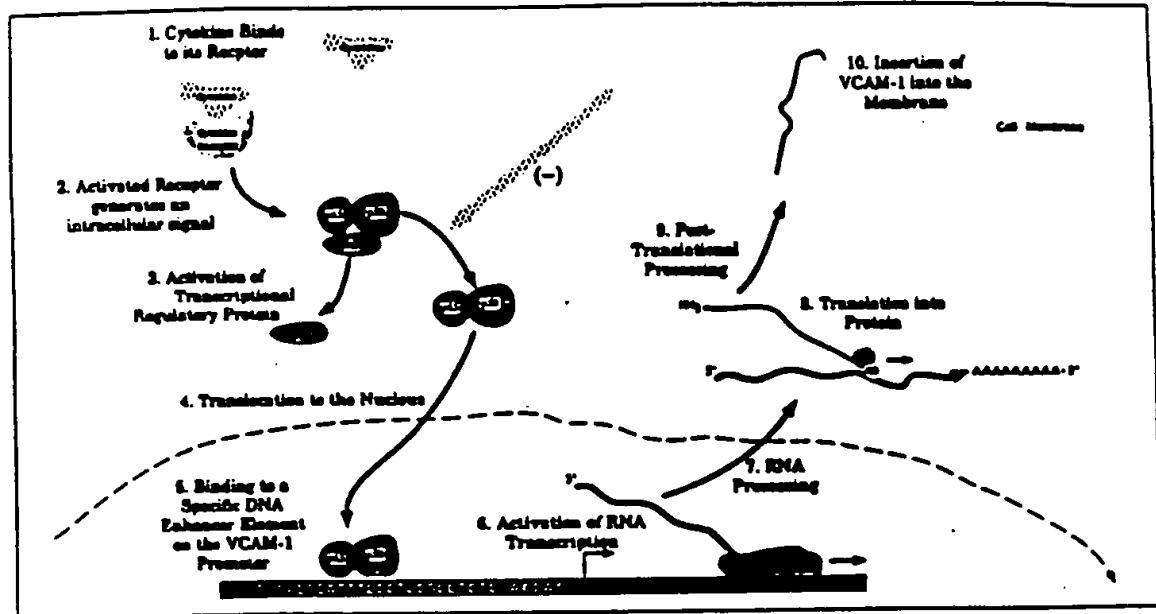
Background of the Invention

This application is in the area of methods and compositions for the treatment of atherosclerosis and other cardiovascular and inflammatory diseases.

Adhesion of leukocytes to the endothelium 5 represents a fundamental, early event in a wide variety of inflammatory conditions, including atherosclerosis, autoimmune disorders and bacterial and viral infections. Leukocyte recruitment to the endothelium is started when inducible adhesion 10 molecule receptors on the surface of endothelial cells interact with counterreceptors on immune cells. Vascular endothelial cells determine which type of leukocytes (monocytes, lymphocytes, or neutrophils) are recruited, by selectively 15 expressing specific adhesion molecules, such as vascular cell adhesion molecule-1 (VCAM-1), intracellular adhesion molecule-1 (ICAM-1), and E-selectin. In the earliest stage of the atherosclerotic lesion, there is a localized 20 endothelial expression of VCAM-1 and selective recruitment of mononuclear leukocytes that express the integrin counterreceptor VLA-4. Because of the selective expression of VLA-4 on monocytes and lymphocytes, but not neutrophils, VCAM-1 is 25 important in mediating the selective adhesion of mononuclear leukocytes. Subsequent conversion leucocytes to foamy macrophages results in the synthesis of a wide variety of inflammatory cytokines, growth factors, and chemoattract 30 that help propagate the leukocyte and plaque recruitment, smooth muscle cell proliferation, endothelial cell activation, and extracellular matrix synthesis characteristic of atherosclerotic plaque.

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VCAM-1 is expressed in cultured human vascular endothelial cells after activation by lipopolysaccharide (LPS) and cytokines such as interleukin-1 (IL-1) and tumor necrosis factor (TNF- $\alpha$ ). These factors are not selective for activation of cell adhesion molecule expression. Scheme 1 illustrates the process of cytokine activation of VCAM-1 gene expression in vascular endothelial cells.



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### Scheme 1

Regulatory schemes for cytokine activation of vascular cell adhesion molecule-1 (VCAM-1) gene expression, through redox sensitive regulatory factors such as NF- $\kappa$ B, in vascular endothelial cells (I $\kappa$ B is an inhibitory subunit; NF- $\kappa$ B is nuclear factor- $\kappa$ B; NH, refers to the amino terminus of the protein and RNA Pol II is RNA polymerase II).

Molecular analysis of the regulatory elements on the human VCAM-1 gene that control its expression suggests an important role for nuclear factor- $\kappa$ B (NF- $\kappa$ B), a transcriptional regulatory factor, or an

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NF- $\kappa$ B like binding protein in oxidation-reduction-sensitive regulation of VCAM-1 gene expression. Transcriptional factors are proteins that activate (or repress) gene expression within the cell

5 nucleus by binding to specific DNA sequences called "enhancer elements" that are generally near the region of the gene, called the "promoter," from which RNA synthesis is initiated. Nuclear factor- $\kappa$ B is a ubiquitously expressed multisubunit

10 transcription factor activated in several cell types by a large and diverse group of inflammatory agents such as TNF $\alpha$ , IL-1 $\beta$ , bacterial endotoxin, and RNA viruses. It plays a key role in mediating inflammatory and other stress signals to the

15 nuclear regulatory apparatus. Although the precise biochemical signals that activate NF- $\kappa$ B are unknown, this transcriptional factor may integrate into a common molecular pathway many of the risk factors and "causative" signals of atherosclerosis, such as hyperlipidemia, smoking, hypertension, and diabetes mellitus.

Importantly, the activation of NF- $\kappa$ B in vascular endothelial cells by diverse signals can be specifically inhibited by antioxidants such as

25 N-acetylcysteine and pyrrolidine dithiocarbamate (see U.S.S.N. 07/969,934, now allowed). This has led to the hypothesis that oxygen radicals play an important role in the activation of NF- $\kappa$ B through an undefined oxidation-reduction mechanism.

30 Because an NF- $\kappa$ B-like enhancer element also regulates the transcription of the VCAM-1 promoter in an oxidation-reduction-sensitive manner, oxidative stress in the atherosclerotic lesion may play a role in regulating VCAM-1 gene expression

35 through this oxidation-reduction-sensitive transcriptional regulatory protein.

It has been hypothesized that modification of low-density lipoprotein (LDL) into oxidatively modified LDL (ox-LDL) by reactive oxygen species is the central event that initiates and propagates 5 atherosclerosis. Steinberg, et al., *N. Engl. J. Med.* 1989; 320:915-924. Oxidized LDL is a complex structure consisting of at least several chemically distinct oxidized materials, each of which, alone or in combination, may modulate cytokine-activated 10 adhesion molecule gene expression. Fatty acid hydroperoxides such as linoleyl hydroperoxide (13-HPODE) are produced from free fatty acids by lipoxygenases and are an important component of oxidized LDL.

15 It has been proposed that a generation of oxidized lipids is formed by the action of the cell lipoxygenase system and that the oxidized lipids are subsequently transferred to LDL. There is thereafter a propagation reaction within the LDL in 20 the medium catalyzed by transition metals and/or sulfhydryl compounds. Previous investigations have demonstrated that fatty acid modification of cultured endothelial cells can alter their susceptibility to oxidant injury. Supplementation 25 of saturated or monounsaturated fatty acids to cultured endothelial cells reduces their susceptibility to oxidant injury, whereas supplementation with polyunsaturated fatty acids (PUFA) enhances susceptibility to oxidant injury.

30 Using reverse-phase HPLC analysis of native and saponified lipid extracts of LDL, it has been demonstrated that 13-HPODE is the predominant oxidized fatty acid in LDL oxidized by activated human monocytes. Chronic exposure to oxidized LDL 35 provides an oxidative signal to vascular endothelial cells, possibly through a specific

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fatty acid hydroperoxide, that selectively augments cytokine-induced VCAM-1 gene expression.

Through a mechanism that is not well defined, areas of vessel wall predisposed to atherosclerosis 5 preferentially sequester circulating LDL. Through a poorly understood pathway, endothelial, smooth muscle, and/or inflammatory cells then convert LDL to ox-LDL. In contrast to LDL, which is taken up through the LDL receptor, monocytes avidly take up 10 ox-LDL through a "scavenger" receptor whose expression, unlike the LDL receptor, is not inhibited as the content of intracellular lipid rises. Thus, monocytes continue to take up ox-LDL and become lipid-engorged macrophage-foam cells 15 that form the fatty streak.

Given that cardiovascular disease is currently the leading cause of death in the United States, and ninety percent of cardiovascular disease is presently diagnosed as atherosclerosis, there is a 20 strong need to identify new methods and pharmaceutical agents for its treatment. Important to this goal is the identification and manipulation of the specific oxidized biological compounds that act as selective regulators of the expression of 25 mediators of the inflammatory process, and in particular, VCAM-1. A more general goal is to identify selective methods for suppressing the expression of redox sensitive genes or activating redox sensitive genes that are suppressed.

30 It is therefore an object of the present invention to provide a treatment for atherosclerosis and other cardiovascular and inflammatory diseases.

35 It is another object of the present invention to provide a method for the selective inhibition of VCAM-1.

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It is still another object of the present invention to provide a method for the treatment of a human disease or disorder that is mediated by the expression or suppression of a redox sensitive 5 gene.

It is another object of the present invention to provide pharmaceutical compositions for the treatment of atherosclerosis and other cardiovascular and inflammatory diseases.

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#### Summary of the Invention

It has been discovered that polyunsaturated fatty acids ("PUFAs") and their hydroperoxides ("ox-PUFAs"), which are important components of 15 oxidatively modified low density lipoprotein (LDL), induce the expression of VCAM-1, but not intracellular adhesion molecule-1 (ICAM-1) or E-selectin in human aortic endothelial cells, through a mechanism that is not mediated by cytokines or other noncytokine signals. This is a fundamental 20 discovery of an important and previously unknown biological pathway in VCAM-1 mediated immune responses.

As nonlimiting examples, linoleic acid, linolenic acid, arachidonic acid, linoleyl 25 hydroperoxide (13-HPODE) and arachidonic hydroperoxide (15-HPETE) induce cell-surface gene expression of VCAM-1 but not ICAM-1 or E-selectin. Saturated fatty acids (such as stearic acid) and monounsaturated fatty acids (such as oleic acid) do 30 not induce the expression of VCAM-1, ICAM-1, or E-selectin.

The induction of VCAM-1 by PUFAs and their fatty acid hydroperoxides is suppressed by the antioxidant pyrrolidine dithiocarbamate (PDTC). 35 This indicates that the induction is mediated by an

oxidized signal molecule, and that the induction is prevented when the oxidation of the molecule is blocked (i.e., the oxidation does not occur), reversed (i.e., the signal molecule is reduced), or 5 when the redox modified signal is otherwise prevented from interacting with its regulatory target.

Cells that are chronically exposed to higher than normal levels of polyunsaturated fatty acids 10 or their oxidized counterparts can initiate an immune response that is not normal and which is out of proportion to the threat presented, leading to a diseased state. The oversensitization of vascular endothelial cells to PUFAS and ox-PUFAS can 15 accelerate the formation, for example, of atherosclerotic plaque.

Based on these discoveries, a method for the treatment of atherosclerosis, post-angioplasty restenosis, coronary artery diseases, angina, and 20 other cardiovascular diseases, as well as noncardiovascular inflammatory diseases that are mediated by VCAM-1, is provided that includes the removal, decrease in the concentration of, or prevention of the formation of oxidized 25 polyunsaturated fatty acids including but not limited to oxidized linoleic ( $C_{18} \Delta^{9,12}$ ), linolenic ( $C_{18} \Delta^{6,9,12}$ ), arachidonic ( $C_{20} \Delta^{5,8,11,14}$ ) and eicosatrienoic ( $C_{20} \Delta^{4,11,14}$ ) acids.

Nonlimiting examples of noncardiovascular 30 inflammatory diseases that are mediated by VCAM-1 include rheumatoid and osteoarthritis, asthma, dermatitis, and multiple sclerosis.

This method represents a significant advance in 35 treating cardiovascular disease, in that it goes beyond the current therapies designed simply to inhibit the progression of the disease, and when used appropriately, provides the possibility to

medically "cure" atherosclerosis by preventing new lesions from developing and causing established lesions to regress.

In an alternative embodiment, a method is provided for suppressing the expression of a redox-sensitive gene or activating a gene that is suppressed through a redox-sensitive pathway, that includes administering an effective amount of a substance that prevents the oxidation of the 10 oxidized signal, and typically, the oxidation of a polyunsaturated fatty acid. Representative redox-sensitive genes that are involved in the presentation of an immune response include, but are not limited to, those expressing cytokines involved 15 in initiating the immune response (e.g., IL-1 $\beta$ ), chemoattractants that promote the migration of inflammatory cells to a point of injury (e.g., MCP-1), growth factors (e.g., IL-6 and the thrombin receptor), and adhesion molecules (e.g., 20 VCAM-1 and E-selectin).

Screens for disorders mediated by VCAM-1 or a redox-sensitive gene are also provided that include the quantification of surrogate markers of the disease. In one embodiment, the level of oxidized 25 polyunsaturated fatty acid, or other appropriate markers, in the tissue or blood, for example, of a host is evaluated as a means of assessing the "oxidative environment" of the host and the host's susceptibility to VCAM-1 or redox-sensitive gene 30 mediated disease.

In another embodiment, the level of circulating or cell-surface VCAM-1 or other appropriate marker and the effect on that level of administration of an appropriate antioxidant is quantified.

35 In yet another assay, the sensitization of a host's vascular endothelial cells to polyunsaturated fatty acids or their oxidized

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counterparts is evaluated. This can be accomplished, for example, by challenging a host with a PUFA or ox-PUFA and comparing the resulting concentration of cell-surface or circulating VCAM-1 or other surrogate marker to a population norm.

5 In another embodiment, in vivo models of atherosclerosis or other heart or inflammatory diseases that are mediated by VCAM-1 can be provided by administering to a host animal an excessive amount of PUFA or oxidized 10 polyunsaturated fatty acid to induce disease. These animals can be used in clinical research to further the understanding of these disorders.

15 In yet another embodiment of the invention, compounds can be assessed for their ability to treat disorders mediated by VCAM-1 on the basis of their ability to inhibit the oxidation of a polyunsaturated fatty acid, or the interaction of a PUFA or ox-PUFA with a protein target.

20 This can be accomplished by challenging a host, for example, a human or an animal such as a mouse, to a high level of PUFA or ox-PUFA and then determining the therapeutic efficacy of a test compound based on its ability to decrease 25 circulating or cell surface VCAM-1 concentration. Alternatively, an in vitro screen can be used that is based on the ability of the test compound to prevent the oxidation of a PUFA, or the interaction of a PUFA or ox-PUFA with a protein target in the 30 presence of an oxidizing substance such as a metal, for example, copper, or an enzyme such as a peroxidase, lipoxygenase, cyclooxygenase, or cytochrome P450.

35 In another embodiment, vascular endothelial cells are exposed to TNF- $\alpha$  or other VCAM-1 inducing material for an appropriate time and then broken by any appropriate means, for example by

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sonication or freeze-thaw. The cytosolic and membrane compartments are isolated. Radiolabeled PUFA is added to defined amounts of the compartments. The ability of the liquid to convert 5 PUFA to ox-PUFA in the presence or absence of a test compound is assayed. Intact cells can be used in place of the broken cell system.

Pyrrolidine dithiocarbamate (PDTC), orally delivered at 25-50mg/kg/day, dramatically inhibited 10 atherogenic fatty streak formation, arterial monocyte-macrophage inflammation, endothelial VCAM-1 expression and essentially normalized endothelium dependent relaxation function in diet induced 15 hypercholesterolemic rabbits with serum cholesterol levels over 1000 mg/dl. At the same doses, other putative therapeutic agents, such as the 20 antioxidants probucol and vitamin E, had no effect on lesion formation in this model.

Endothelial dependent arterial relaxation is 25 restored in experimental atherosclerosis by administration of PDTC. In the diet induced hypercholesterolemic rabbit model, orally delivered (25-50mg/kg/day) PDTC restored endothelial dependent vasoreactivity. This was determined by 30 ring-contraction studies of excised aorta from control and test animals. In patients with atherosclerosis, this manifests itself as normalization of peripheral vascular reactivity in response to hyperemia as measured by non-invasive 35 Doppler flow studies. This is a standardized, commonly available and easily administered test that can be used to titrate functional drug levels to oral doses. The PDTC functions as an anti-ischemic therapy by rapidly normalizing the pathological loss of endothelial derived arterial 40 vasodilation characteristic of cardiovascular diseases and atherosclerosis. This improvement in

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vascular blood flow is manifested as an improvement in symptom and ischemia-limited exercise function and provides a non-invasive assessment of vascular protection. Other clinical indications of 5 abnormalities in endothelial derived vasorelaxation include impotence.

The molecular regulator factory that controls VCAM-1 gene transcription is a novel transcription factor complex consisting of the p65 and p50 10 subunits of the NF- $\kappa$ B/Rel family cross-coupled to the c-fos and the c-jun subunits of the AP-1 family. By both structural and functional studies, it has been established that these AP-1 factors play an important role in the regulation of the 15 VCAM-1 promoter that likely are central to therapeutic regulation of VCAM-1 gene expression. This is the first demonstration of a functional role of this cross-coupled transcription complex in the regulation of an endogenous gene.

20

#### Brief Description of the Figures

Figure 1 is a graph of the cell-surface expression (O.D. 450 nm) of VCAM-1 as a function of hours in human aortic endothelial cells on exposure to the cytokine TNF- $\alpha$  (closed circle); linoleic acid (closed triangle); and linoleyl hydroperoxide (13-HPODE, closed square); and in the absence of 25 exposure to these substances (control, open square).

Figure 2 is a graph of the cell-surface expression (O.D. 450 nm) of VCAM-1 in human aortic 30 endothelial cells on exposure to linoleic acid (closed triangle) and linoleyl hydroperoxide (13-HPODE, closed square) as a function of the concentration of fatty acid ( $\mu$ M).

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Figure 3 is a bar chart graph of the cell-surface expression (O.D. 450 nm) of VCAM-1, ICAM-1 and E-selectin in human aortic endothelial cells on exposure to the cytokine TNF- $\alpha$ , stearic acid, oleic acid, linoleic acid, linolenic acid, and arachidonic acid.

Figure 4 is a bar chart graph of the cell-surface expression (O.D. 450 nm) of VCAM-1 in human aortic endothelial cells on exposure to linoleic acid, 13-HPODE, arachidonic acid, and arachidonic acid hydroperoxide (15-HPETE), with (solid black) or without (hatched lines) the antioxidant pyrrolidine dithiocarbamate.

Figure 5 is an illustration of an autoradiogram indicating the acute induction of VCAM-1 mRNA by linoleic acid and 13-HPODE. HAEC were exposed or not to linoleic acid (7.5  $\mu$ M), 13-HPODE (7.5  $\mu$ M) or TNF- $\alpha$  (100 U/ml). Total RNA was isolated and 20  $\mu$ g was size-fractionated by denaturing 1.0% agarose-formaldehyde gel electrophoresis, transferred to nitrocellulose, and hybridized to either  $^{32}$ P-labeled human A) VCAM-1 specific or B)  $\beta$ -actin specific cDNA. After washing, the filters were exposed to X-ray film at -70°C with one intensifying screen for 24 hours. Identification of lanes: 1) control; 2) linoleic acid (acute, 8-hour exposure); 3) linoleic acid (48-hour exposure); 4) 13-HPODE (acute, 8-hour exposure); and 5) TNF- $\alpha$  (100 U/ml, 4-hour exposure).

Figure 6 is an illustration of an autoradiogram that indicates that induction of VCAM-1 mRNA by polyunsaturated fatty acids is independent of cellular protein synthesis. HAEC were exposed to either linoleic (7.5  $\mu$ M) or arachidonic (7.5  $\mu$ M) acid in the presence or absence of cycloheximide (10  $\mu$ g/ml) for a 4-hour period, and then treated as described in Figure 5.

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Figure 7 is an illustration of an autoradiogram that indicates that linoleic acid induces transcriptional activation of the VCAM-1 promoter by a redox-sensitive NF- $\kappa$ B like factor. HAEC were 5 split at the ratio to give approximately 60% confluence in 100-mm tissue culture plates. HAEC were transfected with either 30  $\mu$ g of p288 VCAMCAT, p85 VCAMCAT, or pSV<sub>2</sub>CAT plasmid by the calcium phosphate coprecipitation technique using standard 10 techniques. After a 24-hour recovery period, HAEC were pretreated or not with 50  $\mu$ M PDTC and after 30 minutes exposed to linoleic acid (7.5  $\mu$ M) or TNF- $\alpha$  (100 U/ml) directly added to the plates. After 18 hours, cell extracts were prepared by rapid 15 freeze-thaw in 0.25 M Tris, pH 8.0. The protein of each cell extract was assayed for chloramphenicol acetyl transferase (CAT) activity, as previously described [Ausubel, 1989] (Ac, acetylated; N, nonacetylated chloramphenicol).

Figure 8 is an illustration of an acrylamide gel slab that indicates that polyunsaturated fatty acids activate NF- $\kappa$ B-like DNA binding activities that are blocked by the antioxidant PDTC. Confluent HAEC in media containing 4% FBS (as described in Figure 1) were pretreated or not with PDTC (50  $\mu$ M) for thirty minutes and then exposed for three hours to linoleic acid (7.5  $\mu$ M, oleic acid (7.5  $\mu$ M), or TNF $\alpha$  (100 U/ml), respectively. Five micrograms of nuclear extract was incubated 25 with a double-stranded <sup>32</sup>P-labeled wtVCAM, size fractionated on 4% native acrylamide gels, and exposed to autoradiography film at -70°C for 18 hours. Two bands A and C, representing NF- $\kappa$ B like binding activity are designated. A weak band B was 30 observed in control (untreated) cells.

Figures 9A and 9B are bar chart graphs of the relative thiobarbituric acid reactive substances

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(O.D. 532 nm) of arachidonic acid and 15-HPETE in the presence or absence of PDTC. The thiobarbituric acid reactivity assay (TBARS) measures the oxidation ability of a material in a 5 cell-free, media-free environment.

Figure 10 is an illustration of an autoradiogram of mRNA, obtained as described below, hybridized to either 32P-labeled human VCAM-1 specific cDNA (Panel 10 A), E-selectin (ELAM-1) specific cDNA (Panel B), or ICAM-1 specific cDNA (Panel C). Following pre-treatment for 30 minutes with 50  $\mu$ M of sodium pyrrolidine dithiocarbamate (PDTC), HUVE (human 15 umbilical vein) cells were exposed to IL-1b (10 U/ml) in the continuous presence of 50  $\mu$ M PDTC. Parallel controls were performed identically except 20 in the absence of PDTC. At the indicated times, total RNA was isolated and 20  $\mu$ g of material size-fractionated by denaturing 1.0% agarose-formaldehyde gel electrophoresis, transferred to 25 nitrocellulose, hybridized as described above, and visualized by autoradiography. Lane 1-0 hour; Lanes 2,4,6,8 - OL-1 alone for 2, 4, 8 and 24 hours, respectively; Lanes 3,5,7,9 - IL-1 with PDTC for 2,4,8 and 24 hours, respectively.

Figure 11 is an illustration of an autoradiogram of mRNA, obtained as described below, hybridized to either 32P-labeled human VCAM-1 specific (Panel A), E-selectin (ELAM-1) specific cDNA (Panel B), or ICAM-1 specific cDNA (Panel C). HUVE cells were 30 pretreated with the indicated concentrations of PDTC, and then exposed to IL-1b in the presence of PDTC for four hours and assayed for VCAM-1 mRNA accumulation by Northern filter hybridization analysis. Lane 1 - control, lane 2 - IL-1 (10u/ml), lane 3 - IL-1b + PDTC (0.05  $\mu$ M), lane 4 - IL-1 LB + PDTC (0.5  $\mu$ M), lane 5 - IL-1b + PDTC (5.0  $\mu$ M), lane 6 - IL=1b + PDTC (50.0  $\mu$ M), lane 7

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- IL-1b + PDTC (100  $\mu$ M).

Figure 12 is an illustration of an autoradiogram of mRNA, obtained as described below, hybridized to either 32P-labeled human VCAM-1 specific cDNA (Panel A), E-selectin (ELAM-1) specific cDNA (Panel B), or ICAM-1 specific cDNA (Panel C). HUVE cells were pretreated as described in Figure 9 with 50  $\mu$ M PDTC, exposed for four hours to the agents indicated below, and assayed for VCAM-1 (Panel A) and ICAM-1 (Panel B) mRNA accumulation. Lane 1 - TNFa (100U/ml), lane 2 - TNFa + PDTC, lane 3 - lipopolysaccharide (LPS) (100ng/ml), lane 4 - LPS + PDTC, lane 5 - poly(I:C) (100mg/ml), lane 6 - poly(I:C) + PDTC.

Figure 13 is a graph of relative cell surface expression of VCAM-1 and ICAM-1 in the presence (dark bars) or absence (white bars) of PDTC and in the presence of multiple types of inducing stimuli. Confluent HUVECs were pretreated or not pretreated (CTL only) for 30 minutes with 50  $\mu$ M PDTC, and then exposed for the indicated times to the indicated agents in the presence or absence (CTL only) of PDTC. Cell surface expression was determined by primary binding with VCAM-1 specific (4B9) and ICAM-1 specific (84H10) mouse monoclonal antibodies followed by secondary binding with a horse-radish peroxidase tagged goat anti-mouse (IgG). Quantitation was performed by determination of calorimetric conversion at 450 nm of TMB. Figure 13 indicates that multiple regulatory signals induce VCAM-1 but not ICAM-1 through a common, dithiocarbamate-sensitive pathway in human vascular endothelial cells.

Figure 14 is a graph of the relative VCAM-1 cell surface expression (O.D. 595 nM) in human umbilical vein endothelial cells, activated by TNFa, versus concentration of various antioxidants. (PDTC is

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sodium N-pyrrolidine dithiocarbamate; DETC is sodium N,N-diethyl-N-carbodithiolate, also referred to as sodium diethyldithiocarbamate; NAC is N-acetyl cysteine; and DF is desferroximine).

5       Figure 15 is a graph of the relative VCAM-1 cell surface expression (O.D. 595 nm) in human umbilical vein endothelial cells, activated by TNF- $\alpha$ , in the presence of the specified amount of antioxidant. (PDTC is sodium N-pyrrolidine dithiocarbamate; DIDTC is sodium N,N-diethyl-N-carbodithioate; SarDTC is sodium N-methyl-N-carboxymethyl-N-carbodithioate; IDADTC is trisodium N,N-di(carboxymethyl)-N-carbodithioate; MGDTC is sodium N-methyl-D-glucamine-N-carbodithioate; MeOBGDTTC is 15 sodium N-(4-methoxybenzyl)-D-glucamine-N-carbodithioate; DEDTC is sodium N,N-diethyl-N-carbodithioate; Di-PDTC is sodium N,N-diisopropyl-N-carbodithioate; NAC is N-acetyl cysteine.)

20       Figure 16 is a graph of the percentage of Molt-4 cells binding to HUVE cells either unstimulated or stimulated with TNF $\alpha$  (100 U/ml) for six hours in the presence or absence of PDTC.

25       Figure 17 is an illustration of the chemical structures of the following active dithiocarbamates: sodium pyrrolidine-N-carbodithioate, sodium N-methyl-N-carboxymethyl-N-carbodithioate, trisodium N,N-di(carboxymethyl)-N-carbodithioate, sodium N-methyl-D-glucamine-N-carbodithioate, sodium N,N-diethyl-N-carbodithioate 30 (sodium diethyldithiocarbamate), and sodium N,N-diisopropyl-N-carbodithioate.

35       Figure 18 is a bar chart graph of the effect of PTDC on the formation of fluorescent adducts of BSA and 13-HPODE, as measured in fluorescent units versus micromolar concentration of PDTC. One micromolar of 13-HPODE was incubated with 200 micrograms of BSA in the presence of PDTC for six

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days. Fluorescence was measured at 430-460 nm with excitation at 330-360 nm.

Figure 19 is a graph of the effect of PTDC on the formation of fluorescent adducts of BSA and ox-PUFA as a function of wavelength (nm) and concentration of PDTC. As the concentration of PDTC increases, the quantity of fluorescent adducts decrease.

Figure 20 is a graph of the effect of PDTC on the oxidation of LDL by horseradish peroxidase (HRP), as measured by the increase in O.D. (234 nm) versus time (minutes) for varying concentrations of PDTC. It is observed that after an incubation period, PDTC inhibits the oxidation of LDL by HRP in a manner that is concentration dependent.

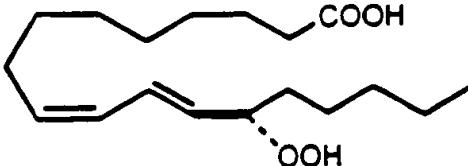
Figure 21 is a chart of the effect of PDTC on the cytokine-induced formation of ox-PUFA in human aortic endothelial cells. As indicated, both TNF- $\alpha$  and IL-1 $\beta$  causes the oxidation of linoleic acid to ox-linoleic acid. The oxidation is significantly prevented by PDTC.

#### Detailed Description of the Invention

##### I. Definitions

As used herein, the term polyunsaturated fatty acid (also referred to herein as a "PUFA") refers to a fatty acid (typically C<sub>8</sub> to C<sub>24</sub>) that has at least two alkenyl bonds, and includes but is not limited to linoleic (C<sub>18</sub>  $\Delta^{9,12}$ ), linolenic (C<sub>18</sub>  $\Delta^{6,9,12}$ ), arachidonic (C<sub>20</sub>  $\Delta^{5,8,11,14}$ ) and eicosatrienoic (C<sub>20</sub>  $\Delta^{8,11,14}$ ) acids.

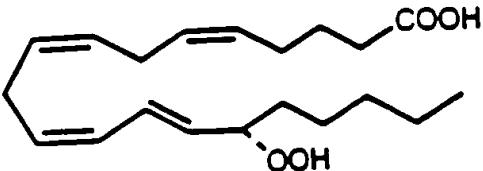
The term oxidized polyunsaturated fatty acid refers to an unsaturated fatty acid in which at least one of the alkenyl bonds has been converted to a hydroperoxide. Nonlimiting examples are:



13-HPODE

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## 15-HPETE



The term alkyl, as used herein, unless otherwise specified, refers to a saturated straight, branched, or cyclic (in the case of C<sub>5</sub> or greater) hydrocarbon of C<sub>1</sub> to C<sub>10</sub> (or lower alkyl, i.e., C<sub>1</sub> to C<sub>3</sub>), which specifically includes methyl, ethyl, propyl, isopropyl, butyl, isobutyl, t-butyl, pentyl, cyclopentyl, isopentyl, neopentyl, hexyl, isohexyl, cyclohexyl, cyclohexylmethyl, 3-methylpentyl, 2,2-dimethylbutyl, and 2,3-dimethylbutyl. The alkyl group can be optionally substituted on any of the carbons with one or more moieties selected from the group consisting of hydroxyl, amino, or mono- or disubstituted amino, wherein the substituent group is independently alkyl, aryl, alkaryl or aralkyl; aryl, alkoxy, aryloxy, nitro, cyano, sulfonic acid, sulfate, phosphonic acid, phosphate, or phosphonate, either unprotected, or protected as necessary, as known to those skilled in the art, for example, as taught in Greene, et al., "Protective Groups in Organic Synthesis," John Wiley and Sons, Second Edition, 1991.

The term alkenyl, as referred to herein, and unless otherwise specified, refers to a straight, branched, or cyclic hydrocarbon of C<sub>2</sub> to C<sub>10</sub> with at least one double bond.

The term alkynyl, as referred to herein, and unless otherwise specified, refers to a C<sub>2</sub> to C<sub>10</sub> straight or branched hydrocarbon with at least one triple bond.

The term aralkyl refers to an aryl group with at least one alkyl substituent.

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The term alkaryl refers to an alkyl group that has at least one aryl substituent.

The term halo (alkyl, alkenyl, or alkynyl) refers to an alkyl, alkenyl, or alkynyl group in which at least one of the hydrogens in the group has been replaced with a halogen atom.

The term aryl, as used herein, and unless otherwise specified, refers to phenyl, biphenyl, or naphthyl, and preferably phenyl. The aryl group can be optionally substituted with one or more moieties selected from the group consisting of alkyl, hydroxyl, amino, alkylamino, arylamino, alkoxy, aryloxy, nitro, cyano, sulfonic acid, sulfate, phosphonic acid, phosphate, or phosphonate,  $\text{CO}_2\text{H}$ , or its pharmaceutically acceptable salt,  $\text{CO}_2(\text{alkyl, aryl, alkaryl or aralkyl})$ , or glucamine, either unprotected, or protected as necessary, as known to those skilled in the art, for example, as taught in Greene, et al., "Protective Groups in Organic Synthesis," John Wiley and Sons, Second Edition, 1991.

The term alkoxy, as used herein, and unless otherwise specified, refers to a moiety of the structure -O-alkyl.

The term acyl, as used herein, refers to a group of the formula  $\text{C}(\text{O})\text{R}'$ , wherein  $\text{R}'$  is an alkyl, aryl, alkaryl or aralkyl group.

The term heteroaryl or heteroaromatic, as used herein, refers to an aromatic moiety that includes at least one sulfur, oxygen, or nitrogen in the aromatic ring. Nonlimiting examples are phenazine, phenothiazine, furyl, pyridyl, pyrimidyl, thienyl, isothiazolyl, imidazolyl, tetrazolyl, pyrazinyl, benzofuranyl, benzothiophenyl, quinolyl, isoquinolyl, benzothienyl, isobenzofuryl, pyrazolyl, indolyl, isoindolyl, benzimidazolyl, purinyl, morpholinyl, carbozolyl, oxazolyl,

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thiazolyl, isothiazolyl, 1,2,4-thiadiazolyl, isooxazolyl, pyrrolyl, pyrazolyl, quinazolinyl, pyridazinyl, pyrazinyl, cinnolinyl, phthalazinyl, quinoxalinyl, xanthinyl, hypoxanthinyl, pteridinyl, 5-azacytidinyl, 5-azauracilyl, triazolopyridinyl, imidazolopyridinyl, pyrrolopyrimidinyl, pyrazolopyrimidinyl, adenine, N<sup>6</sup>-alkylpurines, N<sup>6</sup>-benzylpurine, N<sup>6</sup>-halopurine, N<sup>6</sup>-vinylpurine, N<sup>6</sup>-acetylenic purine, N<sup>6</sup>-acyl purine, N<sup>6</sup>-hydroxyalkyl purine, N<sup>6</sup>-thioalkyl purine, thymine, cytosine, 6-azapyrimidine, 2-mercaptopyrimidine, uracil, N<sup>3</sup>-alkylpyrimidines, N<sup>3</sup>-benzylpyrimidines, N<sup>3</sup>-halopyrimidines, N<sup>3</sup>-vinylpyrimidine, N<sup>3</sup>-acetylenic pyrimidine, N<sup>3</sup>-acyl pyrimidine, N<sup>3</sup>-hydroxyalkyl purine, and N<sup>3</sup>-thioalkyl purine, and isoxazolyl. The heteroaromatic group can be optionally substituted as described above for aryl. The heteroaromatic can be partially or totally hydrogenated as desired. As a nonlimiting example, dihydropyridine can be used in place of pyridine. Functional oxygen and nitrogen groups on the heterocyclic base can be protected as necessary or desired during the reaction sequence. Suitable protecting groups are well known to those skilled in the art, and include trimethylsilyl, dimethylhexylsilyl, t-butyldimethylsilyl, and t-butyldiphenylsilyl, tritylmethyl, alkyl groups, acyl groups such as acetyl and propionyl, methylsulfonyl, and p-toluylsulfonyl.

The term hydroxyalkyl, as used herein, refers to a C<sub>1</sub> to C<sub>6</sub> alkyl group in which at least one of the hydrogens attached to any of the carbon atoms is replaced with a hydroxy group.

The term thiol antioxidant refers to a sulfur containing compound that retards oxidation.

The term pharmaceutically acceptable derivative refers to a derivative of the active compound that

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upon administration to the recipient, is capable of providing directly or indirectly, the parent compound, or that exhibits activity itself.

The term "pharmaceutically acceptable cation" 5 refers to an organic or inorganic moiety that carries a positive charge and that can be administered in association with a pharmaceutical agent, for example, as a counterion in a salt. Pharmaceutically acceptable cations are known to 10 those of skill in the art, and include but are not limited to sodium, potassium, and quaternary amine.

The term "physiologically cleavable leaving group" refers to a moiety that can be cleaved in vivo from the molecule to which it is attached, and 15 includes but is not limited to an organic or inorganic anion, a pharmaceutically acceptable cation, acyl (including but not limited to (alkyl)C(O), including acetyl, propionyl, and butyryl), alkyl, phosphate, sulfate and sulfonate.

20 The term "enantiomerically enriched composition or compound" refers to a composition or compound that includes at least 95%, and preferably at least 97, 98, 99, or 100% by weight of a single enantiomer of the compound.

25 The term amino acid includes synthetic and naturally occurring amino acids, including but not limited to, for example, alanyl, valinyl, leucinyl, isoleucinyl, prolinyl, phenylalaninyl, tryptophanyl, methioninyl, glycyl, serinyl, 30 threoninyl, cysteinyl, tyrosinyl, asparaginyl, glutaminyl, aspartoyl, glutaoyl, lysinyl, arginyl, and histidinyl.

A "linking moiety" as used herein, is any divalent group that links two chemical residues, 35 including but not limited to alkyl, alkenyl, alkynyl, aryl, polyalkyleneoxy (for example -[(CH<sub>2</sub>)<sub>n</sub>O-]<sub>n</sub>-, -C<sub>1-6</sub>alkoxy-C<sub>1-10</sub>alkyl-,

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-C<sub>1-6</sub>alkylthio-C<sub>1-10</sub> alkyl-, -NR<sup>3-</sup>, and -(CHOH)<sub>n</sub>CH<sub>2</sub>OH,  
wherein n is independently 0, 1, 2, 3, 4, 5, or 6.

5           II.       Identification of Oxidized and Unoxidized  
                  Polyunsaturated Fatty Acids as Direct  
                  Mediators of VCAM-1 Expression

To establish whether a PUFA or oxidized PUFA  
acts as a direct immunomodulator of endothelial  
cell gene expression, early passaged human aortic  
endothelial cells (HAEC) were cultured for eight  
10 hours in media and serum and exposed to saturated  
(stearic), monounsaturated (oleic), and  
polyunsaturated (linoleic and arachidonic) fatty  
acids; as well as with the fatty acid  
hydroperoxides of linoleic (13-HPODE) or  
15 arachidonic (15-HPETE) acids. HAEC were also  
alternatively exposed to the cytokine tumor  
necrosis factor- $\alpha$ .

HAEC were exposed to linoleic acid or 13-HPODE  
for varying times up to 48 hours and then assayed  
20 for cell surface VCAM-1 expression by ELISA assay.  
The results were compared to HAEC exposed to the  
cytokine TNF- $\alpha$  (100 U/ml) for the same time  
periods. VCAM-1 expression in HAEC incubated with  
either linoleic acid or 13-HPODE is transiently  
25 induced. The expression peaks at approximately 8-9  
hours with significant expression at 24 hours and  
then decreases by 48 hours. The kinetics of VCAM-1  
induction by both linoleic acid and 13-HPODE mirror  
that of TNF- $\alpha$ , and thus the mechanisms by which  
30 polyunsaturated fatty acids induce VCAM-1 thus  
appear to be similar to that of TNF- $\alpha$ .

35 Dose-response studies of linoleic acid and  
13-HPODE on VCAM-1 gene expression at 8 hours were  
also conducted. It was observed that 7.5  $\mu$ M is  
the lowest peak dose by which linoleic acid and

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13-HPODE induces significant VCAM-1 gene expression.

It was then explored whether short term incubation of endothelial cells with 5 polyunsaturated fatty acids induces ICAM-1 and E-selectin expression as well. It was determined that the polyunsaturated fatty acids linoleic and arachidonic acids induced cell-surface gene expression to approximately 59% of TNF-induced gene 10 expression of VCAM-1. Strikingly, neither ICAM-1 nor E-selectin were induced by these fatty acids. Conversely, the saturated fatty acid stearic acid and the monounsaturated fatty acid oleic acid did not induce the expression of VCAM-1, ICAM-1, or 15 E-selectin. VCAM-1 gene expression was also observed by incubation of HAEc with the oxidized metabolites of linoleic acid (13-HPODE) and arachidonic acid (15-HPETE).

To investigate whether oxidative stress in 20 endothelial cells provided by polyunsaturated fatty acids and their oxidized metabolites induces VCAM-1 through a redox-sensitive mechanism, HAEc were pretreated with the antioxidant pyrrolidine dithiocarbamate (PDTC, 50  $\mu$ M) for 30 minutes and 25 then the cells were independently incubated with linoleic acid, arachidonic acid, 13-HPODE, and 15-HPETE (all 7.5  $\mu$ M) for 8 hours. It was determined that PDTC suppressed the gene expression 30 of VCAM-1 induced by the polyunsaturated fatty acids and their oxidized counterparts. This indicates that the induction is mediated by a oxidized signal molecule, and that the induction is prevented when the oxidation of the molecule is 35 blocked (i.e., the oxidation does not occur), reversed (i.e., the signal molecule is reduced), or its interaction with a target protein prevented, perhaps through a redox complex.

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To determine whether the selective induction of VCAM-1 by PUFAs and their oxidized metabolites is observed at the mRNA level, HAEc were incubated with linoleic acid or 13-HPODE. Linoleic acid and 5 13-HPODE induced VCAM-1 mRNA accumulation that was similar to levels induced by TNF- $\alpha$ . In contrast, there was no induction of ICAM-1 or E-selectin gene expression at the mRNA level in HAEc incubated with linoleic acid or 13-HPODE. The findings mimic 10 those found at the cell-surface level. These results indicate that pretranslational regulatory mechanisms mediate induction of VCAM-1 gene expression by polyunsaturated fatty acids and their oxidative metabolites.

15 It was also desired to determine whether polyunsaturated fatty acids work as a primary signal or operate through a regulatory protein involving the cytokine IL-4 in inducing VCAM-1 gene expression. To investigate whether newly 20 synthesized proteins such as IL-4 are involved in the synthesis and gene expression of VCAM-1 induced by PUFAs such as linoleic acid, HAEc were incubated with 13-HPODE (7.5  $\mu$ M) and exposed to the protein synthesis inhibitor, cycloheximide. There was no 25 inhibition of mRNA accumulation of VCAM-1 by cycloheximide in HAEc incubated with 13-HPODE. The production of IL-4 by HAEc incubated with linoleic or arachidonic acids and their oxidative metabolites, as determined by ELISA was also 30 measured. There was no increase in IL-4 output by HAEc incubated with these PUFAs or their oxidized metabolites.

35 Previous investigations have demonstrated through deletion and heterologous promoter studies that cytokines and non-cytokines activate VCAM-1 gene expression in endothelial cells at least in part transcriptionally through two NF- $\kappa$ B-like DNA

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binding elements. It has also been demonstrated that PDTC inhibits VCAM-1 gene expression through a redox-sensitive NF- $\kappa$ B like factor. To determine whether polyunsaturated fatty acids induce 5 transcriptional activation of the human VCAM-1 promoter via a similar mechanism, the chimeric reporter gene p288 VCAM-CAT, containing coordinates -288 to +22 of the human VCAM-1 promoter, was transiently transfected into HAEC. The addition of 10 linoleic acid (7.5  $\mu$ M) induced VCAM-1 promoter. The addition of linoleic acid (7.5  $\mu$ M) induced VCAM-1 promoter activity that was over two fold that of the control and approximately 60% of the maximum signal induced by TNF- $\alpha$ . Similar results 15 were obtained with the minimal cytokine-inducible promoter of the VCAM-1 gene (p85 VCAM-CAT), containing the -77 and -63 bp NF- $\kappa$ B-like sites. Neither linoleic acid nor TNF- $\alpha$  had any effect on activity using a constitutively expressed pSV<sub>2</sub> CAT 20 construct. PDTC inhibited the transcriptional activation of both VCAM-1 promoter constructs induced by linoleic acid. The data indicate that analogous to TNF- $\alpha$ , polyunsaturated fatty acids such as linoleic acid induce the transcriptional 25 activation of VCAM-1 through an NF- $\kappa$ B-like redox-sensitive mechanism.

To determine whether polyunsaturated fatty acids and their oxidative metabolites regulate VCAM-1 promoter activity through an NF- $\kappa$ B-like 30 transcriptional regulatory factor, nuclear extracts from HAEC were assayed for DNA binding activity to a double-stranded oligonucleotide containing the VCAM-1 NF- $\kappa$ B-like promoter elements located at positions -77 and -63. As shown in Figure 7, two 35 bands A and C, representing NF- $\kappa$ B-like activity were induced in response to a three hour exposure to linoleic acid (7.5  $\mu$ M). Similar findings were

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observed on exposure to the cytokine TNF- $\alpha$  (100 U/ml). A weak band B was observed in control (untreated) cells. No induction of NF- $\kappa$ B-like binding was observed with the monounsaturated fatty acid oleic acid. Pretreatment of the cells for thirty minutes with PDTC inhibited the A and C complex DNA binding activity after linoleic acid activation. These findings are similar to previously reported findings that PDTC blocks the activation of VCAM-1 gene expression in HUVEC by inhibiting the activation of these NF  $\kappa$ B-like DNA binding proteins.

15 Example 1 Effect of Oxidized and Unoxidized Polyunsaturated Fatty Acids on the Kinetics of the Activation of VCAM-1 Gene Expression

Human aortic endothelial cells (HAEC) were plated in 96 well plates and incubated with linoleic acid (7.5  $\mu$ M), 13-HPODE (7.5  $\mu$ M), or TNF- $\alpha$  (100 U/ml) at five different time points up to 48 hours. HAEC, obtained from Clonetics (Boston, MA), were cultured in Medium 199 supplemented with 20% fetal bovine serum (FBS), 16 U/ml heparin, 10 U/ml epidermal growth factor, 50  $\mu$ g/ml endothelial cell growth supplement, 2 mM L-glutamine, 100 U/ml penicillin, and 100  $\mu$ g/ml streptomycin. One day before the experiment, cells were placed in a medium containing 4% FBS. Confluent HAEC were incubated for up to 48 hours with TNF- $\alpha$  (100 U/ml), or stearic, oleic, linoleic, linolenic, or arachidonic acids (7.5  $\mu$ M). Similar studies were performed with differing doses of linoleic acid or 13-HPODE for an 8 hour period (1-60  $\mu$ M) (Figure 2). Quantitation was performed by determination of colorimetric conversion at 450 nm of TMB. Studies were performed in triplicate (n=4).

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for each experimental value). \*-value differs (p<0.05) from Control.

As shown in Figure 1, both linoleic acid and 13-HPODE induced the expression of VCAM-1. At ten hours after exposure, the amount of cell surface VCAM-1 induced by linoleic acid and 13-HPODE was greater than half that induced by the cytokine TNF- $\alpha$ .

As shown in Figure 2, the induction of VCAM-1 by linoleic acid and 13-HPODE is concentration sensitive. At a concentration of between 2 and 10  $\mu$ M of these compounds, there is a sharp increase in the amount of induced cell surface VCAM-1, which then remains approximately constant up to a concentration of at least 100  $\mu$ M. It should be observed that the PUFA concentration indicated in Figure 2 is in addition to that found endogenously in HAEC.

20      **Example 2      Polyunsaturated Fatty Acids Induce Gene Expression of VCAM-1 but not ICAM-1 or E-selectin.**

The cell surface expression of VCAM-1, ICAM-1, and E-selectin was measured in HAEC by ELISA. 25 HAEC, obtained from Clonetics (California), were cultured in Medium 199 supplemented with 20% fetal bovine serum (FBS), 16 U/ml heparin, 10 U/ml epidermal growth factor, 50  $\mu$ g/ml endothelial cell growth supplement, 2 mM L-glutamine, 100 U/ml penicillin, and 100  $\mu$ g/ml streptomycin. One day 30 before the experiment, cells were placed in a medium containing 4% FBS. Confluent HAEC were incubated or not for 8 hours with TNF- $\alpha$  (100 U/ml), or stearic, oleic, linoleic, linolenic, or 35 arachidonic acids (7.5  $\mu$ M). Cell-surface expression of A) VCAM-1, B) ICAM-1, and C) E-selectin was determined by primary binding with VCAM-1 specific, ICAM-1 specific, and E-selectin

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specific mouse antibodies followed by secondary binding with a horseradish peroxidase-tagged goat anti-mouse (IgG). Quantitation was performed by determination of colorimetric conversion at 450 nm of TMB. Studies were performed in triplicate (n=4 for each experimental value). \*-value differs (p<0.05) from Control.

As shown in Figure 3, linoleic acid, linolenic acid, and arachidonic acid significantly induced the expression of VCAM-1, but did not induce the cell-surface expression of ICAM-1 or E-selectin. Neither stearic acid nor oleic acid induced the expression of VCAM-1, ICAM-1, or E-selectin. TNF- $\alpha$  strongly induced the expression of all three cell-surface molecules.

**Example 3 The Antioxidant PDTC Suppresses VCAM-1 Induction by Polyunsaturated Fatty Acids and their Oxidative Metabolites.**

Confluent HAEC were pretreated in the presence or absence of PDTC (sodium pyrrolidine dithiocarbamate, 50  $\mu$ M) for thirty minutes. The cells were then incubated for eight hours with TNF- $\alpha$  (100 U/ml), linoleic or arachidonic acid (7.5  $\mu$ M), or the fatty acid hydroperoxides 13-HPODE (7.5  $\mu$ M) or 15-HPETE (7.5  $\mu$ M). The cell surface expression of VCAM-1 was measured in HAEC by ELISA, as described in Example 1. Studies were performed in triplicate (n = 4 for each experimental value). \*-value differs (p<0.05) from control.

As indicated in Figure 4, PDTC suppresses the induction of VCAM-1 by linoleic acid, 13-HPODE, arachidonic acid and 15-HPETE.

**Example 4 Acute Induction of VCAM-1 mRNA by Linoleic Acid and 13-HPODE.**

HAEC were exposed to linoleic acid (7.5  $\mu$ M) or 13-HPODE (7.5  $\mu$ M). Total RNA was isolated and 20

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μg size-fractionated by denaturing 1.0% agarose-formaldehyde gel electrophoresis, transferred to nitrocellulose, and hybridized to either <sup>32</sup>P-labeled human A) VCAM-1 specific or B) 5  $\beta$ -actin specific cDNA and visualized by autoradiography. After washes, filters were exposed to X-ray film at -70°C with one intensifying screen for 24 hours. Identification of lanes: 1) control; 2) linoleic acid (acute, 10 8-hour exposure); 3) linoleic (48-hour exposure); 4) 13-HPODE (acute, 8-hour exposure); and 5) TNF- $\alpha$  (100 U/ml, 4-hour exposure).

As shown in Figure 5, both linoleic acid and 13-HPODE induce the production of mRNA for VCAM-1 in 15 eight hours. After 48 hours, linoleic acid no longer induces VCAM-1 mRNA.

**Example 5 Induction of VCAM-1 mRNA by PUFAs is Independent of Cellular Protein Synthesis.**

20 HAEC were exposed to either linoleic or arachidonic acid (7.5  $\mu$ M) in the presence or absence of cycloheximide (10  $\mu$ g/ml) for a 4-hour period. Total RNA was isolated and 20  $\mu$ g was size-fractionated by denaturing 1.0% agarose-formaldehyde gel electrophoresis, 25 transferred to nitrocellulose, and hybridized to A) <sup>32</sup>P-labeled human VCAM-1 or B)  $\beta$ -actin specific cDNA and then visualized by autoradiography. After washes, filters were exposed to X-ray film at -70°C with one intensifying screen for 24 hours.

30 As indicated in Figure 6, the induction of VCAM-1 by linoleic and arachidonic acids are independent of cellular protein synthesis.

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**Example 6** Linoleic acid induces transcriptional activation of the VCAM-1 promoter by a redox-sensitive NF- $\kappa$ B like factor.

HAEC were split at the ratio to give approximately 60% confluence in 100-mm tissue culture plates. HAEC were transfected with either 30  $\mu$ g of p288 VCAMCAT, p85 VCAMCAT, or pSV<sub>2</sub>CAT plasmid by the calcium phosphate coprecipitation technique using standard techniques. After a 24-hour recovery period, HAEC were pretreated with 50  $\mu$ M PDTC and after 30 minutes exposed to linoleic acid (7.5  $\mu$ M) or TNF- $\alpha$  (100 U/ml) directly added to the plates. After 18 hours, cell extracts were prepared by rapid freeze-thaw in 0.25 M Tris, pH 8.0. Protein of each cell extract was assayed for chloramphenicol acetyl transferase (CAT) activity (Ac, acetylated; N, nonacetylated chloramphenicol).

Figure 7 illustrates the results of this experiment. Linoleic acid induces transcriptional activation of the VCAM-1 promoter by a redox-sensitive NF- $\kappa$ B like factor. These results are similar to those observed by the activation of VCAM-1 promotor by cytokines such as TNF- $\alpha$ . This suggests that PUFAs act through an oxidized intermediate that also mediates the cytokine activation of VCAM-1.

**Example 7** Polyunsaturated Fatty Acids Activate NF- $\kappa$ B-like DNA Binding Activities that are Blocked by the Antioxidant PDTC.

Confluent HAEC in media containing 4% FBS (as described in Example 1) were pretreated with PDTC (50  $\mu$ M) for 30 minutes and then exposed for 3 hours to linoleic acid or oleic acid (7.5  $\mu$ M), or TNF- $\alpha$  (100 U/ml). Five micrograms of nuclear extract was incubated with a double-stranded <sup>32</sup>P-labeled wtVCAM, size fractionated on 4% native acrylamide gels, and exposed to autoradiography film at -70°C for 18

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hours. Two bands A and C, representing NF- $\kappa$ B like binding activity are designated. A weak band B was observed in control (untreated) cells.

Figure 8 illustrates that linoleic acid induces 5 NF- $\kappa$ B binding activity to VCAM-1 promotor in a redox-sensitive manner. This is analogous to cytokine TNF- $\alpha$  and suggests a similar mechanism of action. TNF- $\alpha$  probably induces VCAM-1 through a mechanism that is mediated by an ox-PUFA.

10 Example 8 Oxidation in a cell-free, media-free setup, by both unoxidized and oxidized (15-HPETE) arachidonic acid

Figures 9A and 9B are bar chart graphs of the relative thiobarbituric acid reactive substances 15 (O.D. 532 nm) of arachidonic acid and 15-HPETE in the presence or absence of PDTC. The thiobarbituric acid reactivity assay (TBARS) measures the oxidation ability of a material in a cell-free, media-free environment. As indicated in 20 the Figures, both arachidonic acid and 15-HPETE showed significant TBARS activity that was inhibited by PDTC.

### III. Method for the Treatment of VCAM-1 Mediated Disorders

25 The discovery that polyunsaturated fatty acids and their oxidized metabolites are selective, redox-sensitive immunomodulators provides a basis for the therapy of disorders that are mediated by VCAM-1 or by redox-sensitive genes.

30 A method for the treatment of atherosclerosis, post-angioplasty restenosis, coronary artery diseases, angina, and other cardiovascular diseases, as well as noncardiovascular inflammatory diseases that are mediated by VCAM-1 is provided 35 that includes the removal, decrease in the

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concentration of, or prevention of the formation of oxidized polyunsaturated fatty acids, including but not limited to oxidized linoleic, linolenic, and arachidonic acids. In an alternative embodiment, a 5 method for the treatment of these diseases is provided that includes the prevention of the interaction of a PUFA or ox-PUFA with a protein or peptide that mediates VCAM-1 expression.

Inhibition of the expression of VCAM-1 can be 10 accomplished in a number of ways, including through the administration of an antioxidant that prevent the oxidation of a polyunsaturated fatty acid, by in vivo modification of the metabolism of PUFAs into ox-PUFAs, as described in more detail below.

15 1. Administration of Antioxidants

Any compound that reduces an ox-PUFA or which 20 inhibits the oxidation of PUFA, and which is relatively nontoxic and bioavailable or which can be modified to render it bioavailable, can be used in this therapy. One of ordinary skill in the art can easily determine whether a compound reduces an ox-PUFA or inhibits the oxidation of PUFA using standard techniques.

**Dithiocarboxylate Antioxidants**

25 It has been discovered that dithiocarboxylates are useful in the treatment of atherosclerosis and other cardiovascular and inflammatory diseases. Dithiocarboxylates, including dithiocarbamates, can be used to block the ability of cells, including 30 endothelial cells, to express VCAM-1 or to suppress the expression of a redox-sensitive gene or activate a gene that is suppressed through a redox-sensitive pathway.

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At least one of the compounds, pyrrolidine dithiocarbamate (PDTC), inhibits VCAM-1 gene expression at a concentration of less than 1.0 micromolar. This compounds also exhibits 5 preferential toxicity to proliferating or abnormally dividing vascular smooth muscle cells. Another dithiocarbamate, sodium N-methyl-N-carboxymethyl-N-carbodithioate, also inhibits the expression of VCAM-1, without significant effect on 10 ICAM-1, but does not exhibit preferential toxicity to abnormally dividing vascular smooth muscle cells. Another dithiocarbamate, sodium N-methyl-N-carboxymethyl-N-carbodithioate, also inhibits the expression of VCAM-1, without significant effect on 15 ICAM-1, but does not exhibit preferential toxicity to abnormally dividing vascular smooth muscle cells.

It has been discovered that pyrrolidine dithiocarbamate does not significantly block ELAM-1 or ICAM-1 expression, and therefore treatment with 20 this compound does not adversely affect aspects of the inflammatory response mediated by ELAM-1 or ICAM-1. Thus, a generalized immunosuppression is avoided. This may avoid systemic complications 25 from generalized inhibition of adhesion molecules in the many other cell types known to express them. Other pharmaceutically acceptable salts of PDTC are also effective agents for the treatment of cardiovascular and inflammatory disorders.

30 Dithiocarbamates are transition metal chelators clinically used for heavy metal intoxication. Baselt, R.C., F.W.J. Sunderman, et al. (1977), "Comparisons of antidotal efficacy of sodium diethyldithiocarbamate, D-penicillamine and 35 triethylenetetramine upon acute toxicity of nickel carbonyl in rats." Res Commun Chem Pathol Pharmacol 18(4): 677-88; Menne, T. and K. Kaaber

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Dithiocarbamates have also been used 25 adjunctively in cis-platinum chemotherapy to prevent renal toxicity. Hacker, M.P., W.B. Ershler, et al. (1982). "Effect of disulfiram (tetraethylthiuram disulfide) and 30 diethyldithiocarbamate on the bladder toxicity and antitumor activity of cyclophosphamide in mice." Cancer Res 42(11): 4490-4. Bodenner, 1986 #733; 35 Saran, M. and Bors, W. (1990). "Radical reactions in vivo--an overview." Radiat. Environ. Biophys. 29(4): 249-62.

-35-

A dithiocarbamate currently used in the treatment of alcohol abuse is disulfiram, a dimer of diethyldithiocarbamate. Disulfuram inhibits hepatic aldehyde dehydrogenase. Inoue, K., and Fukunaga, et al., (1982). "Effect of disulfiram and its reduced metabolite, diethyldithiocarbamate on aldehyde dehydrogenase of human erythrocytes." Life Sci 30(5): 419-24.

It has been reported that dithiocarbamates inhibit HIV virus replication, and also enhance the maturation of specific T cell subpopulations. This has led to clinical trials of diethyldithiocarbamate in AIDS patient populations. Reisinger, E., et al., (1990). "Inhibition of HIV progression by dithiocarb." Lancet 335: 679.

Dithiocarboxylates are compounds of the structure A-SC(S)-B, which are members of the general class of compounds known as thiol antioxidants, and are alternatively referred to as carbodithiols or carbodithiolates. It appears that the -SC(S)- moiety is essential for therapeutic activity, and that A and B can be any group that does not adversely affect the efficacy or toxicity of the compound.

In an alternative embodiment, one or both of the sulfur atoms in the dithiocarbamate is replaced with a selenium atom. The substitution of sulfur for selenium may decrease the toxicity of the molecule in certain cases, and may thus be better tolerated by the patient.

A and B can be selected by one of ordinary skill in the art to impart desired characteristics to the compound, including size, charge, toxicity, and degree of stability, (including stability in an acidic environment such as the stomach, or basic environment such as the intestinal tract). The selection of A and B will also have an important

-36-

effect on the tissue-distribution and pharmacokinetics of the compound. In general, for treatment of cardiovascular disease, it is desirable that the compound accumulate, or 5 localize, in the arterial intimal layer containing the vascular endothelial cells. The compounds are preferably eliminated by renal excretion.

An advantage in administering a dithio-carboxylate pharmaceutically is that it does not 10 appear to be cleaved enzymatically in vivo by thioesterases, and thus may exhibit a prolonged halflife in vivo.

In a preferred embodiment, A is hydrogen or a pharmaceutically acceptable cation, including but 15 not limited to sodium, potassium, calcium, magnesium, aluminum, zinc, bismuth, barium, copper, cobalt, nickel, or cadmium; a salt-forming organic acid, typically a carboxylic acid, including but not limited to acetic acid, oxalic acid, tartaric acid, succinic acid, malic acid, ascorbic acid, 20 benzoic acid, tannic acid, pamoic acid, alginic acid, polyglutamic acid, naphthalenesulfonic acid, naphthalenedisulfonic acid, or polygalacturonic acid; or a cation formed from ammonia or other 25 nitrogenous base, including but not limited to a nitrogenous heterocycle, or a moiety of the formula  $NR^4R^5R^6R^7$ , wherein R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, and R<sup>7</sup> are independently hydrogen, C<sub>1-6</sub> linear, branched, or (in the case of C<sub>4-6</sub>) cyclic alkyl, hydroxy(C<sub>1-6</sub>)alkyl (wherein one or 30 more hydroxyl groups are located on any of the carbon atoms), or aryl, N,N-dibenzyl-ethylenediamine, D-glucosamine, choline, tetraethylammonium, or ethylenediamine.

In another embodiment, A can be a 35 physiologically cleavable leaving group that can be cleaved in vivo from the molecule to which it is attached, and includes but is not limited acyl

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(including acetyl, propionyl, and butyryl), alkyl, phosphate, sulfate or sulfonate.

In one embodiment, B is alkyl, alkenyl, alkynyl, alkaryl, aralkyl, haloalkyl, haloalkenyl,

5 haloalkynyl, aryl, alkaryl, hydrogen, C<sub>1-4</sub> alkoxy-C<sub>1-10</sub> alkyl, C<sub>1-4</sub> alkylthio-C<sub>1-10</sub> alkyl, NR<sup>2</sup>R<sup>3</sup>, -(CHOH)<sub>n</sub>CH<sub>2</sub>OH, wherein n is 0, 1, 2, 3, 4, 5, or 6, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>1</sup>, including alkylacetyl, alkylpropionyl, and alkylbutyryl, or hydroxy(C<sub>1-4</sub>)alkyl- (wherein one or 10 more hydroxyl groups are located on any of the carbon atoms).

In another embodiment, B is NR<sup>2</sup>R<sup>3</sup>, wherein R<sup>2</sup> and R<sup>3</sup> are independently alkyl; -(CHOH)<sub>n</sub>(CH<sub>2</sub>)<sub>n</sub>OH, wherein n is 0, 1, 2, 3, 4, 5, or 6; -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>1</sup>,

15 -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>4</sup>; hydroxy(C<sub>1-4</sub>)alkyl-; alkenyl (including but not limited to vinyl, allyl, and CH<sub>3</sub>CH=CH-CH<sub>2</sub>CH<sub>3</sub>); alkyl(CO<sub>2</sub>H), alkenyl(CO<sub>2</sub>H), alkynyl(CO<sub>2</sub>H), or aryl, wherein the aryl group can be substituted as described above, notably, for 20 example, with a NO<sub>2</sub>, CH<sub>3</sub>, t-butyl, CO<sub>2</sub>H, halo, or p-OH group; or R<sup>2</sup> and R<sup>3</sup> can together constitute a bridge such as -(CH<sub>2</sub>)<sub>m</sub>- , wherein m is 3, 4, 5, or 6, and wherein R<sup>4</sup> is alkyl, aryl, alkaryl, or aralkyl, including acetyl, propionyl, and butyryl.

25 In yet another embodiment, B can be a heterocyclic or alkylheterocyclic group. The heterocycle can be optionally partially or totally hydrogenated. Nonlimiting examples are those listed above, including phenazine, phenothiazine, 30 pyridine and dihydropyridine.

In still another embodiment, B is the residue of a pharmaceutically-active compound or drug. The term drug, as used herein, refers to any substance used internally or externally as a medicine for the treatment, cure, or prevention of a disease or 35 disorder.

Nonlimiting examples are drugs for the treatment or prevention of cardiovascular disease, including 5 antioxidants such as probucol; nicotinic acid; agents that prevent platelets from sticking, such as aspirin; antithrombotic agents such as coumadin; calcium channel blockers such as varapamil, diltiazem, and nifedipine; angiotensin converting enzyme (ACE) inhibitors such as captopril and enalopril,  $\beta$ -blockers such as propanalol, 10 terbutalol, and labetalol, nonsteroidal antiinflammatories such as ibuprofen, indomethacin, fenoprofen, mefenamic acid, flufenamic acid, sulindac, or corticosteroids. The  $-C(S)SA$  group can be directly attached to the drug, or attached 15 through any suitable linking moiety.

In another embodiment, the dithiocarbamate is an amino acid derivative of the structure  $AO_2C-R^9-NR^{10}-C(S)SA$ , wherein  $R_9$  is a divalent  $B$  moiety, a linking moiety, or the internal residue of any of the 20 naturally occurring amino acids (for example,  $CH_2CH$  for alanine,  $CH_3$  for glycine,  $CH(CH_3)_2NH_2$  for lysine, etc.), and  $R^{10}$  is hydrogen or lower alkyl.

$B$  can also be a polymer to which one or more dithiocarbamate groups are attached, either 25 directly, or through any suitable linking moiety. The dithiocarbamate is preferably released from the polymer under in vivo conditions over a suitable time period to provide a therapeutic benefit. In a preferred embodiment, the polymer itself is also 30 degradable in vivo. The term biodegradable or bioerodible, as used herein, refers to a polymer that dissolves or degrades within a period that is acceptable in the desired application (usually in vivo therapy), usually less than five years, and 35 preferably less than one year, on exposure to a physiological solution of pH 6-8 having a temperature of between 25 and 37°C. In a preferred

embodiment, the polymer degrades in a period of between 1 hour and several weeks, according to the application.

A number of degradable polymers are known.

5 Nonlimiting examples are peptides, proteins, nucleoproteins, lipoproteins, glycoproteins, synthetic and natural polypeptides and polyamino acids, including but not limited to polymers and copolymers of lysine, arginine, asparagine, 10 aspartic acid, cysteine, cystine, glutamic acid, glutamine, hydroxylysine, serine, threonine, and tyrosine; polyorthoesters, including poly( $\alpha$ -hydroxy acids), for example, polylactic acid, polyglycolic acid, poly(lactide-co-glycolide), polyanhydrides, 15 albumin or collagen, a polysaccharide containing sugar units such as lactose, and polycaprolactone. The polymer can be a random or block copolymer.

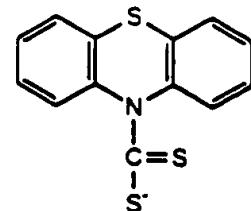
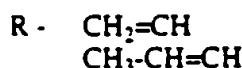
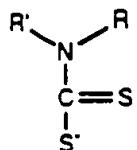
B can also be a group that enhances the water solubility of the dithiocarbamate, for example, 20 -lower alkyl-O-R<sup>8</sup>, wherein R<sup>8</sup> is -PO<sub>2</sub>(OH)M<sup>+</sup> or PO<sub>3</sub>(M<sup>+</sup>), wherein M<sup>+</sup> is a pharmaceutically acceptable cation; -C(O)(CH<sub>2</sub>)<sub>2</sub>CO<sub>2</sub>M<sup>+</sup>, or -SO<sub>3</sub>M<sup>+</sup>; -lower alkylcarbonyl-lower alkyl; -carboxy lower alkyl; -lower alkylamino-lower alkyl; N,N-di-substituted amino 25 lower alkyl-, wherein the substituents each independently represent lower alkyl; pyridyl-lower alkyl-; imidazolyl-lower alkyl-; imidazolyl-Y-lower alkyl wherein Y is thio or amino; morpholinyl-lower alkyl; pyrrolidinyl-lower alkyl; thiazolinyl-lower 30 alkyl-; piperidinyl-lower alkyl; morpholinyl-lower hydroxylalkyl; N-pyrryl; piperazinyl-lower alkyl; N-substituted piperazinyl-lower alkyl, wherein the substituent is lower alkyl; triazolyl-lower alkyl; tetrazolyl-lower alkyl; tetrazolylamino-lower 35 alkyl; or thiazolyl-lower alkyl.

In an alternative embodiment, a dimer such as B-C(S)S-SC(S)-B can be administered.

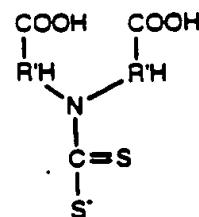
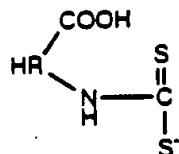
-40-

Nonlimiting examples of dithiocarbamates are those of the structure:

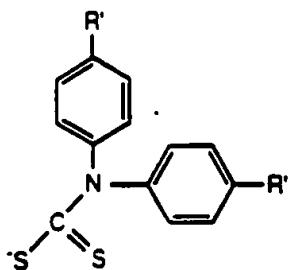
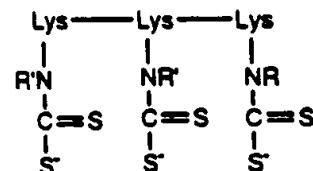
## 1) Aliphatic Substrate



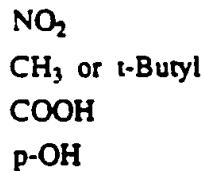
## 2) Amino Acid



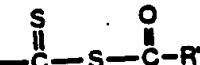
## Polyamino acid



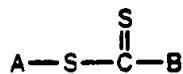
## Substituents



R -

Na<sup>+</sup>

Thioester

Ca<sup>++</sup>NH<sub>4</sub><sup>+</sup>Choline<sup>+</sup> and quaternary aminesMg<sup>++</sup>Al<sup>+++</sup>K<sup>+</sup>H<sup>+</sup>

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Dithiocarboxylates should be chosen for use in treating atherosclerosis and other cardiovascular and inflammatory diseases that have the proper lipophilicity to locate at the affected site. The 5 compound should not compartmentalize in low turnover regions such as fat deposits. In a preferred embodiment for treatment of cardiovascular disease, the pharmacokinetics of the compound should not be dramatically affected by 10 congestive heart failure or renal insufficiency.

For topical applications for the treatment of inflammatory skin disorders, the selected compound should be formulated to be absorbed by the skin in a sufficient amount to render a therapeutic effect 15 to the afflicted site.

The dithiocarboxylate must be physiologically acceptable. In general, compounds with a therapeutic index of at least 2, and preferably at least 5 or 10, are acceptable. The therapeutic 20 index is defined as the  $EC_{50}/IC_{50}$ , wherein  $EC_{50}$  is the concentration of compound that inhibits the expression of VCAM-1 by 50% and  $IC_{50}$  is the concentration of compound that is toxic to 50% of the target cells. Cellular toxicity can be 25 measured by direct cell counts, trypan blue exclusion, or various metabolic activity studies such as  $^3H$ -thymidine incorporation, as known to those skilled in the art. The therapeutic index of PDTc in tissue culture is over 100 as measured by 30 cell toxicity divided by ability to inhibit VCAM-1 expression activated by TNF $\alpha$ , in HUVE cells. Initial studies on the rapidly dividing cell type HT-18 human glioma demonstrate no toxicity at concentrations 100-fold greater than the 35 therapeutic concentration. Disulfiram, an orally administered form of diethyldithiocarbamate, used in the treatment of alcohol abuse, generally

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elicits no major clinical toxicities when administered appropriately.

There are a few dithiocarbamates that are known to be genotoxic. These compounds do not fall 5 within the scope of the present invention, which is limited to the administration of physiologically acceptable materials. An example of a genotoxic dithiocarbamate is the fungicide zinc dimethyldithiocarbamate. Further, the 10 anticholinesterase properties of certain dithiocarbamates can lead to neurotoxic effects. Miller, D. (1982). Neurotoxicity of the pesticidal carbamates. Neurobehav. Toxicol. Teratol. 4(6): 779-87.

15 The term dithiocarboxylate as used herein specifically includes, but is not limited to, dithiocarbamates of the formulas:



wherein R<sup>1</sup> is H or a pharmaceutically acceptable 20 cation, including but not limited to sodium, potassium, or NR<sup>4</sup>R<sup>5</sup>R<sup>6</sup>R<sup>7</sup>, wherein R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup>, and R<sup>7</sup> are independently hydrogen, C<sub>1-6</sub> linear, branched, or cyclic alkyl, hydroxy(C<sub>1-6</sub>)alkyl (wherein one or more hydroxyl groups are located on any of the carbon 25 atoms), or aryl, and

R<sup>2</sup> and R<sup>3</sup> are independently C<sub>1-10</sub> linear, branched or cyclic alkyl; -(CHOH)<sub>n</sub>(CH<sub>2</sub>)<sub>n</sub>OH, wherein n is 0, 1, 2, 3, 4, 5, or 6; -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>4</sup>, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>4</sup>; hydroxy(C<sub>1-6</sub>)alkyl-, or R<sup>2</sup> and R<sup>3</sup> together constitute 30 a bridge such as -(CH<sub>2</sub>)<sub>m</sub>-, wherein m is 3-6, and wherein R<sup>4</sup> is alkyl, aryl, alkaryl, or aralkyl, including acetyl, propionyl, and butyryl.

Specific examples of useful dithiocarbamates, 35 illustrated in Figure 15, include sodium pyrrolidine-N-carbodithioate, sodium N-methyl-N-carboxymethyl-N-carbodithioate, trisodium N,N-di(carboxymethyl)-N-carbodithioate, sodium N-

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methyl-D-glucamine-N-carbodithioate, sodium N,N-diethyl-N-carbodithioate (sodium diethyldithiocarbamate), and sodium N,N-diisopropyl-N-carbodithioate.

5 The active dithiocarboxylates and in particular, dithiocarbamates are either commercially available or can be prepared using known methods.

## II. Biological Activity

10 The ability of dithiocarboxylates to inhibit the expression of VCAM-1 can be measured in a variety of ways, including by the methods set out in detail below in Examples 9 to 15. For convenience, Examples 9-11 and 14-15 describe the evaluation of the biological activity of sodium pyrrolidine-N-15 carbodithioate (also referred to as PDTC). These examples are not intended to limit the scope of the invention, which specifically includes the use of any of the above-described compounds to treat atherosclerosis, and other types of inflammation 20 and cardiovascular disease mediated by VCAM-1. Any of the compounds described above can be easily substituted for PDTC and evaluated in similar fashion.

25 Examples 12 and 13 provide comparative data on the ability of a number of dithiocarbamates to inhibit the gene expression of VCAM-1. The examples below establish that the claimed dithiocarbamates specifically block the ability of VCAM-1 to be expressed by vascular endothelial 30 cells in response to many signals known to be active in atherosclerosis and the inflammatory response.

### Experimental Procedures

35 Cell Cultures HUVE cells were isolated from human umbilical veins that were cannulated,

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perfused with Hanks solution to remove blood, and then incubated with 1% collagenase for 15 minutes at 37°C. After removal of collagenase, cells were cultured in M199 medium supplemented with 20% fetal bovine serum (HyClone), 16 µg/ml heparin (ESI Pharmaceuticals, Cherry Hill, NJ), 50 µg/ml endothelial cell growth supplement (Collaborative Research Incorporated, Bedford MA), 25 mM Hepes Buffer, 2 mM L-glutamin, 100 µg/ml penicillin and 100 µg/ml streptomycin and grown at 37°C on tissue culture plates coated 0.1% gelatin. Cells were passaged at confluence by splitting 1:4. Cells were used within the first 8 passages.

Incubation with Cytokines and Other Reagents

Confluent HUVE cells were washed with phosphate buffered saline and then received fresh media. The indicated concentrations of PDTC were added as pretreatment 30 minutes before adding cytokines. Cytokines and other inducers were directly added to medium for the times and at the concentrations indicated in each experiment. Human recombinant IL-1b was the generous gift of Upjohn Company (Kalamazoo, Michigan). TNFa was obtained from Boehringer Engelheim. Bacterial lipopolysaccharide (LPS), polyinosinic acid: polycytidilic acid (Poly I:C), and pyrrolidine dithiocarbamate (PDTC) were obtained from Sigma Chemical (St. Louis, MO). All other reagents were of reagent grade.

RNA Isolation: Total cellular RNA was isolated by a single extraction using an acid guanidium thiocyanate-phenol-chloroform mixture. Cells were rinsed with phosphate buffered saline and then lysed with 2 ml of guanidium isothiocyanate. The solution was acidified with 0.2 ml of sodium acetate (pH 4.0) and then extracted with 2 ml phenol and 0.4 ml chloroform:isoamyl alcohol (24:1). The RNA underwent two ethanol

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precipitations prior to being used for Northern blot analysis.

Northern Blot Analysis: Total cellular RNA (20 µg) was size fractionated using 1% agarose 5 formaldehyde gels in the presence of 1 ug/ml ethidium bromide. The RNA was transferred to a nitrocellulose filter and covalently linked by ultraviolet irradiation using a Stratlinker UV crosslinker (Stratagene, La Jolla, CA). 10 Hybridizations were performed at 42°C for 18 hours in 5X SSC (1X=150 mM NaCl, 15 mM Na citrate), 1% sodium dodecyl sulfate, 5X Denhardt solution, 50% formamide, 10% dextran sulfate and 100 ug/ml of sheared denatured salmon sperm DNA. Approximately 15 1-2X 10<sup>6</sup> cpm/ml of labeled probe (specific activity > 108 cpm/ug DNA) were used per hybridization. Following hybridization, filters were washed with a final stringency of 0.2X SSC at 55°C. The nitrocellulose was stripped using boiled water 20 prior to rehybridization with other probes. Autoradiography was performed with an intensifying screen at -70°C.

<sup>32</sup>P Probes: <sup>32</sup>P labeled DNA probes were made using the random primer oligonucleotide method. The 25 ICAM-1 probe was an Eco R1 fragment of human cDNA. The ELAM-1 probe was a 1.85 kb Hind III fragment of human cDNA. The VCAM-1 probe was a Hind III-Xho I fragment of the human cDNA consisting of nucleotide 132 to 1814.

30 Enzyme Linked Immunosorbent Assay (ELISA): HUVE cells were plated on 96-well tissue culture plates 48 to 72 hours before the assay. Primary antibodies in M199 with 5% FBS were added to each well and incubated one hour at 37°C. The cells 35 were then washed and incubated for one hour with peroxidase conjugated goat anti-mouse IgG (Bio Rad) diluted 1/500 in M199 with 5% FBS. The wells were

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then washed and binding of antibody was detected by the addition of 100  $\mu$ l of 10 mg/ml 3,3,5,5'-tetramethyl-benzidine (Sigma) with 0.003% H<sub>2</sub>O<sub>2</sub>. The reaction was stopped by the addition of 25  $\mu$ l of 8N 5 sulfuric acid. Plates were read on an ELISA reader (Bio Rad) at OD 450 nm after blanking on rows stained only with second step antibody. Data represent the means of triplicate.

Antibodies: Monoclonal antibody (MAb) 4B9 10 recognizing vascular cell adhesion molecule-1 (VCAM-1) was the generous gift of Dr. John Harlan (University of Washington). MAb E9A1F1 recognizing endothelial cell adhesion molecule (ELAM-1) was the generous gift of Dr. Swerlick (Emory University). 15 Hybridomas producing mAb 84H10 recognizing intercellular adhesion molecule 1 (ICAM-1) are routinely grown in our laboratory and antibody was used as tissue culture supernatant.

20 **Example 9      PDTC Blocks IL-1b Mediated Induction of HUVEC VCAM-1, but not ICAM-1 or ELAM-1, mRNA Accumulation**

To determine whether the oxidative state of the endothelial cell can alter the basal or induced expression of cell adhesion molecule genes, 25 cultured human vascular endothelial cells were exposed to the inducing cytokine, IL-1b (10 U/ml) in the presence or absence of the thiolated metal chelating antioxidant, pyrrolidine dithiocarbamate (PDTC, 50  $\mu$ M) for up to 24 hours. As shown in 30 Figure 10, IL-1b alone (lanes 2, 4, 6, 8) induces the expected rapid and transient induction of VCAM-1 (Panel A), E-selectin (ELAM-1, Panel B) and ICAM-1 (Panel C) mRNA accumulation, all of which peak at four hours. However, in the presence of PDTC, IL-35 1b induction of VCAM-1 mRNA accumulation is dramatically inhibited by over 90% (panel A, lanes 3, 5, 7, 9). In contrast, although IL-1b mediated

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induction of ELAM-1 is slightly inhibited at 2 and 24 hours (compare lane 2 and 3, 8 and 9, panel B), PDTc does not inhibit the induction at 4 and 8 hours (lane 5 and 7, panel B). IL-1b mediated induction of ICAM-1 mRNA accumulation is not affected (panel B, lanes 3, 5, 7, 9). Indeed, a mild augmentation of IL-1b induction of ICAM-1 mRNA accumulation (~30%) is observed (compare lanes 4 and 5, panel B). Equivalent amounts of nitrocellulose transferred RNA per lane was confirmed by ethidium bromide staining and visualization.

A dose-response analysis was performed to determine whether PDTc inhibits the induction of VCAM-1 gene expression by IL-1b in a dose dependent manner. As shown in Figure 11, PDTc inhibits IL-1b mediated induction of VCAM-1 gene expression with a steep dose-response curve (Figure 11, panel A) with a calculated EC<sub>50</sub> of approximately 10  $\mu$ M, while PDTc does not inhibit IL-1b mediated induction of ELAM-1 expression with these concentrations (Fig. 11, panel B). The IL-1b mediated induction of ICAM-1 mRNA accumulation is enhanced by PDTc with the concentration higher than 0.5  $\mu$ M (Fig. 2, compare lane 2 and lane 4-7, panel C).

These data demonstrate that IL-1b utilizes a dithiocarboxylate, and in particular, a dithiocarbamate sensitive step as part of its signaling mechanism in the induction of VCAM-1 gene expression. The data also appear to indicate that this dithiocarbamate sensitive step does not play a significant role in the IL-1b mediated induction of ELAM-1 or ICAM-1 gene expression.

Example 10 PDTc Blocks Induction of HUVEC VCAM-1 mRNA Accumulation by Multiple Stimuli

To determine whether other well-described activators of VCAM-1 gene expression also utilize a

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PDTC sensitive step, three distinct classes of activators were tested: another classic receptor mediated inducing agent (TNFa), a non-receptor mediated inducer (lipopolysaccharide (LPS)) and a 5 recently described novel inducer (double stranded RNA, poly(I:C)). In all three cases, PDTC dramatically inhibited the induction of VCAM-1 mRNA accumulation in HUVECs after four hours (Figure 12, Panel A). Although the TNFa mediated ELAM-1 gene 10 expression is suppressed to some extent (Fig. 12 lane 1 and 2, panel B), LPS and poly(I:C) mediated ELAM-1 mRNA accumulation was unaffected (Fig. 12 lane 3-6, panel B). The induction of ICAM-1 mRNA accumulation was unaffected (Figure 12, Panel C). 15 This data indicates that structurally distinct inducing agents, acting through distinct pathways, share a common regulatory step specific for the induction of VCAM-1 gene expression.

20 **Example 11 PDTC Blocks HUVE Cell Surface Expression of VCAM-1 Induced by Multiple Stimuli**

To determine whether, like its mRNA, the induction of endothelial cell surface protein expression of VCAM-1 could also be inhibited by 25 PDTC, monoclonal antibodies were used in an ELISA assay to quantitate the induction of cell surface VCAM-1 and ICAM-1 in cultured HUVE cells. As shown in Figure 13, multiple classes of activating agents, in the absence of PDTC (-PDTC), induce the 30 rapid and transient accumulation of VCAM-1 (top left panel) at the cell surface peaking at six hours. In the presence of PDTC (+PDTC, top right panel), the induction of cell surface expression of VCAM-1 by all agents tested is dramatically 35 inhibited (80-90%). In contrast, the induced expression of cell surface ICAM-1 is unaffected

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under identical conditions (bottom left and right panels).

These data demonstrate that, like its mRNA accumulation, cell surface VCAM-1 expression are  
5 selectively inhibited by dithiocarbamates and that multiple classes of activating agents utilize a similar, dithiocarbamate sensitive mechanism to induce VCAM-1 gene expression.

10 **Example 12 Comparative Effectiveness of  
Antioxidants in Blocking TNFa  
Induction of VCAM-1**

To determine whether structurally similar or dissimilar antioxidants could also inhibit VCAM-1 gene expression, and with what potency, HUVE cells  
15 were exposed to TNFa for six hours in the presence or absence of different concentrations of four different antioxidants. As shown in Figure 14, both diethyldithiocarbamate (DETC) and N-acetyl cysteine (NAC) inhibited VCAM-1 expression at  
20 concentrations of 5  $\mu$ M and 30  $\mu$ M, respectively. In contrast, PDTC (PDTC) was effective between 5 and 50  $\mu$ M. The iron metal chelator, desferroximine, had no effect at the concentrations tested.

25 **Example 13 PDTC Inhibits TNF Induction of VCAM-1/VLA-4 Mediated Adhesion**

The ability of a variety of antioxidants to inhibit TNF- $\alpha$  induction of VCAM-1 in HUVE cells was evaluated by the method set out in Example 12. Figure 15 is a graph of the relative VCAM-1 cell surface expression (O.D. 595 nM) in TNF- $\alpha$  activated HUVE cells versus concentrations of PTDC (sodium N-pyrrolidine dithiocarbamate), DIDTC (sodium N,N-diethyl-N-carbodithioate), SarDTC (sodium N-methyl-N-carboxymethyl-N-carbodithioate), IDADTC  
30 (trisodium N,N-di(carboxymethyl)-N-carbodithioate),  
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MGDTC (sodium N-methyl-D-glucamine-N-carbodithioate), MeOBGDTc (sodium N-(4-methoxybenzyl)-D-glucamine-N-carbodithioate), DEDTC (sodium N,N-diethyl-N-carbodithioate), Di-PDTC (sodium N,N-diisopropyl-N-carbodithioate), and NAC is (N-acetyl cysteine).

5                   **Example 13    PDTC Inhibits TNF Induction of VCAM-1/VLA-4 Mediated Adhesion**

10                  In order to define whether PDTC inhibition of VCAM-1 regulation is associated with functional consequences, the binding of Molt-4 cells to HUVEC cells either unstimulated or stimulated with TNFa (100U/ml) was examined for six hours in the presence or absence of PDTC. Molt-4 cells have 15 been previously shown to bind to activated HUVEC via a VCAM-1 dependent mechanism. As shown in Figure 16, the percentage of Molt-4 binding to HUVEC cells decreased when PDTC was present in the media.

20                  **Example 14    PDTC Inhibits Monocyte Binding to the Thoracic Aorta of Cholesterol Fed Rabbits**

25                  An experiment was performed to determine whether the thiol antioxidant PDTC would be efficacious in blocking the first monocyte binding component of atherosclerosis in an experimental animal model. One mature New Zealand white rabbit (3.5 Kg) received an intravenous injection of PDTC (20 mg/Kg, as a concentration of 20 mg/ml in PBS) once 30 daily for 5 days. Injections were given via an indwelling cannula in the marginal ear vein, which was kept patent by flushing with heparinized saline solution. The PDTC solution was mixed fresh daily or on alternate days (stored light-protected at 35 4°C), and filtered (0.45 mm pore filter) just prior to use. After the first injection, when the

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cannula was placed, the drug was administered with the rabbit in the conscious state without apparent discomfort or other ill effect. On the second day of injections, the rabbit was given chow containing 5 1% cholesterol by weight, which was continued throughout the remainder of the experiment. On the 10 fifth day, the animal was euthanized and the thoracic aorta was excised and fixed. After appropriate preparation, the sample was imaged on 15 the lower stage of an ISI DS-130 scanning electron microscope equipped with a LaB emitter. Using dual-screen imaging and a transparent grid on the CRT screen, 64 adjacent fields at a 620x magnification were assessed, to cover an area of 15 -1.3 mm<sup>2</sup>. Within each field, the number of adherent leukocytes (WBC) and erythrocytes (RBC) were counted and recorded.

The data from the arch sample are as follows: 5 WBC and ~25 RBC per 1.3 mm<sup>2</sup> field. This level of 20 WBC adhesion is similar to control animals fed regular chow (about 7 per field have been seen in arch and thoracic samples from 2 'negative control' experiments). 'Positive control' rabbits fed 1% cholesterol for 4 days but not given antioxidant 25 show about a 5-fold increase in adhesion, to 38 WBC/1.3 mm<sup>2</sup>. A considerable amount of mostly cell-sized debris was observed adherent to each arch sample. It is unclear whether this material is an artifact of preparation, or was present *in vivo*, 30 and if so, whether it is related to PDTC administration. These studies suggest that PDTC infusions can effectively block initial monocyte adhesion to the aortic endothelium.

35 **Example 15 Inhibition of BSA 13-HPODE Adducts with PDTC**  
Figure 18 is a bar chart graph of the effect of PDTC on the formation of fluorescent adducts of BSA

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and 13-HPODE, as measured in fluorescent units versus micromolar concentration of PDTC. One micromolar of 13-HPODE was incubated with 200 micrograms of BSA in the presence of PDTC for six 5 days. Fluorescence was measured at 430-460 nm with excitation at 330-360 nm. For details of the assay, see Freebis, J., Parthasarathy, S., Steinberg, D, Proceedings of the National Academy of Sciences 89, 10588-10592, 1992. In a typical 10 reaction 100 nmols of LOOH (generated by the lipoxygenase catalyzed oxidation of linoleic acid) in incubated with 100  $\mu$ g of bovine serum albumin for 48 to 72 hours and the formation of fluorescent 15 products are followed by measuring the fluorescent spectrum with excitation at 360 nm and emission between 390 and 500 nm.

As indicated, PDTC decreases the concentration of fluorescent adducts of BSA and 13-HPODE.

Figure 19 is a graph of the effect of PTDC on 20 the formation of fluorescent adducts of BSA and ox-PUFA as a function of wavelength (nm) and concentration of PDTC. As the concentration of PDTC increases, the quantity of fluorescent adducts decrease.

25 **Example 16 Effect of PDTC on the oxidation of LDL by horseradish peroxidase**

Figure 20 is a graph of the effect of PDTC on the oxidation of LDL by horseradish peroxidase (HRP), as measured over time (minutes) for varying 30 concentrations of PDTC. The oxidation of LDL was followed by measuring the oxidation of the fatty acid components of LDL as determined by the increase in optical density at 234 nm. When a polyunsaturated fatty acid is oxidized, there is a 35 shift of double bonds resulting in the formation of conjugated dienes which absorb at 234 nm. The

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intercept of the initiation and propagation curve (lag phase) is suggested to be a measure of the oxidizability of LDL. Higher the lag phase, more resistant is the LDL to oxidation. Typically 100 5  $\mu$ g of human LDL is incubated with 5  $\mu$ M  $H_2O_2$  and the increase in absorption of 234 nm is followed.

It is observed that after an incubation period, PDTC inhibits the oxidation of LDL by HRP in a manner that is concentration dependent.

10 **Example 17 Effect of PDTC on the cytokine-induced formation of ox-PUFA**

Figure 21 is a chart of the effect of PDTC on the cytokine-induced formation of ox-PUFA in human aortic endothelial cells. As indicated, both TNF- $\alpha$  15 and IL-1B causes the oxidation of linoleic acid to ox-linoleic acid. The oxidation is significantly prevented by PDTC.

2. Modification of the Synthesis and Metabolism of PUFAs and ox-PUFAs

20 Inhibition of the expression of VCAM-1 can be accomplished via a modification of the metabolism of PUFAs into ox-PUFAs. For example, a number of enzymes are known to oxidize unsaturated materials, including peroxidases, lipoxygenases, 25 cyclooxygenases, and cytochrome P450. The inhibition of these enzymes may prevent the oxidation of PUFAs in vivo. PUFAs can also be oxidized by metal-dependent nonenzymatic materials.

30 **IV. Method for Modifying the Expression of a Redox-Sensitive Gene**

In an alternative embodiment, a method is provided for suppressing the expression of a redox-sensitive gene or activating a gene that is

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suppressed through a redox-sensitive pathway, that includes administering an effective amount of a substance that prevents the oxidation of the oxidized signal, and typically, the oxidation of a 5 polyunsaturated fatty acid. Representative redox-sensitive genes that are involved in the presentation of an immune response include, but are not limited to, those expressing cytokines involved in initiating the immune response (e.g., IL-1 $\beta$ ), 10 chemoattractants that promote the migration of inflammatory cells to a point of injury (e.g., MCP-1), growth factors (IL-6, thrombin receptor), and adhesion molecules (e.g., VCAM-1 and E-selectin).

15 Given this disclosure, one of ordinary skill in the art will be able to screen a wide variety of antioxidants for their ability to suppress the expression of a redox-sensitive gene or activate a gene that is suppressed through a redox-sensitive 20 pathway. All of these embodiments are intended to fall within the scope of the present invention.

Based on the results of this screening, nucleic acid molecules containing the 5' regulatory sequences of the redox-sensitive genes can be used 25 to regulate or inhibit gene expression *in vivo* can be identified. Vectors, including both plasmid and eukaryotic viral vectors, may be used to express a particular recombinant 5' flanking region-gene construct in cells depending on the preference and 30 judgment of the skilled practitioner (see, e.g., Sambrook et al., Chapter 16). Furthermore, a number of viral and nonviral vectors are being developed that enable the introduction of nucleic acid sequences *in vivo* (see, e.g., Mulligan, 1993 35 Science, 260, 926-932; United States Patent No. 4,980,286; United States Patent No. 4,868,116; incorporated herein by reference). Recently, a

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delivery system was developed in which nucleic acid is encapsulated in cationic liposomes which can be injected intravenously into a mammal. This system has been used to introduce DNA into the cells of 5 multiple tissues of adult mice, including endothelium and bone marrow (see, e.g., Zhu et al., 1993 Science 261, 209-211; incorporated herein by reference).

The 5' flanking sequences of the redox-sensitive 10 gene can be used to inhibit the expression of the redox-sensitive gene. For example, an antisense RNA of all or a portion of the 5' flanking region of the redox-sensitive gene can be used to inhibit expression of the gene *in vivo*. Expression vectors 15 (e.g., retroviral expression vectors) are already available in the art which can be used to generate an antisense RNA of a selected DNA sequence which is expressed in a cell (see, e.g., U.S. Patent No. 4,868,116; U.S. Patent No. 4,980,286).

20 Accordingly, DNA containing all or a portion of the sequence of the 5' flanking region of the gene can be inserted into an appropriate expression vector so that upon passage into the cell, the transcription of the inserted DNA yields an 25 antisense RNA that is complementary to the mRNA transcript of the gene normally found in the cell. This antisense RNA transcript of the inserted DNA can then base-pair with the normal mRNA transcript found in the cell and thereby prevent the mRNA from 30 being translated. It is of course necessary to select sequences of the 5' flanking region that are downstream from the transcriptional start sites for the redox-sensitive gene to ensure that the antisense RNA contains complementary sequences 35 present on the mRNA. Antisense RNA can be generated *in vitro* also, and then inserted into cells. Oligonucleotides can be synthesized on an

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automated synthesizer (e.g., Model 8700 automated synthesizer of Milligen-Bioscience, Burlington, MA or ABI Model 380B). In addition, antisense deoxyoligonucleotides have been shown to be 5 effective in inhibiting gene transcription and viral replication (see e.g., Zamecnik et al., 1978 Proc. Natl. Acad. Sci. USA 75, 280-284; Zamecnik et al., 1986 Proc. Natl. Acad. Sci., 83, 4143-4146; Wickstrom et al., 1988 Proc. Natl. Acad. Sci. USA 85, 1028-1032; Crooke, 1993 FASEB J. 7, 533-539. Furthermore, recent work has shown that improved inhibition of expression of a gene by antisense oligonucleotides is possible if the antisense oligonucleotides contain modified nucleotides (see, 10 15 e.g., Offensperger et. al., 1993 EMBO J. 12, 1257-1262 (in vivo inhibition of duck hepatitis B viral replication and gene expression by antisense phosphorothioate oligodeoxynucleotides); Rosenberg et al., PCT WO 93/01286 (synthesis of sulfurthioate 20 25 oligonucleotides); Agrawal et al., 1988 Proc. Natl. Acad. Sci. USA 85, 7079-7083 (synthesis of antisense oligonucleoside phosphoramidates and phosphorothioates to inhibit replication of human immunodeficiency virus-1); Sarin et al., 1989 Proc. Natl. Acad. Sci. USA 85, 7448-7794 (synthesis of antisense methylphosphonate oligonucleotides); Shaw et al., 1991 Nucleic Acids Res 19, 747-750 (synthesis of 3' exonuclease-resistant oligonucleotides containing 3' terminal 30 35 phosphoroamidate modifications); incorporated herein by reference).

The sequences of the 5' flanking region of the redox-sensitive gene can also be used in triple helix (triplex) gene therapy. Oligonucleotides complementary to gene promoter sequences on one of the strands of the DNA have been shown to bind promoter and regulatory sequences to form local

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triple nucleic acid helices which block transcription of the gene (see, e.g., 1989 Maher et al., Science 245, 725-730; Orson et al., 1991 Nucl. Acids Res. 19, 3435-3441; Postal et al., 1991 Proc. Natl. Acad. Sci. USA 88, 8227-8231; Cooney et al., 1988 Science 241, 456-459; Young et al., 1991 Proc. Natl. Acad. Sci. USA 88, 10023-10026; Duval-Valentin et al., 1992 Proc. Natl. Acad. Sci. USA 89, 504-508; 1992 Blume et al., Nucl. Acids Res. 20, 1777-1784; 1992 Grigoriev et al., J. Biol. Chem. 267, 3389-3395.

Recently, both theoretical calculations and empirical findings have been reported which provide guidance for the design of oligonucleotides for use in oligonucleotide-directed triple helix formation to inhibit gene expression. For example, oligonucleotides should generally be greater than 14 nucleotides in length to ensure target sequence specificity (see, e.g., Maher et al., (1989); Grigoriev et al., (1992)). Also, many cells avidly take up oligonucleotides that are less than 50 nucleotides in length (see e.g., Orson et al., (1991); Holt et al., 1988 Mol. Cell. Biol. 8, 963-973; Wickstrom et al., 1988 Proc. Natl. Acad. Sci. USA 85, 1028-1032). To reduce susceptibility to intracellular degradation, for example by 3' exonucleases, a free amine can be introduced to a 3' terminal hydroxyl group of oligonucleotides without loss of sequence binding specificity (Orson et al., 1991). Furthermore, more stable triplexes are formed if any cytosines that may be present in the oligonucleotide are methylated, and also if an intercalating agent, such as an acridine derivative, is covalently attached to a 5' terminal phosphate (e.g., via a pentamethylene bridge); again without loss of sequence specificity (Maher et al., (1989); Grigoriev et al., (1992)).

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Methods to produce or synthesize oligo-nucleotides are well known in the art. Such methods can range from standard enzymatic digestion followed by nucleotide fragment isolation (see 5 e.g., Sambrook et al., Chapters 5, 6) to purely synthetic methods, for example, by the cyanoethyl phosphoramidite method using a Milligen or Beckman System 1Plus DNA synthesizer (see also, Ikuta et al., in Ann. Rev. Biochem. 1984 53, 323-356 10 (phosphotriester and phosphite-triester methods); Narang et al., in Methods Enzymol., 65, 610-620 (1980) (phosphotriester method)). Accordingly, DNA sequences of the 5' flanking region of the redox-sensitive gene described herein can be used to 15 design and construct oligonucleotides including a DNA sequence consisting essentially of at least 15 consecutive nucleotides, with or without base modifications or intercalating agent derivatives, for use in forming triple helices specifically 20 within the 5' flanking region of a redox-sensitive gene in order to inhibit expression of the gene.

In some cases it may be advantageous to insert 25 enhancers or multiple copies of the regulatory sequences into an expression system to facilitate screening of methods and reagents for manipulation of expression.

#### V. Models and Screens

Screens for disorders mediated by VCAM-1 or a redox-sensitive gene are also provided that include 30 the quantification of surrogate markers of the disease. In one embodiment, the level of oxidized polyunsaturated fatty acid, or other appropriate markers, in the tissue or blood, for example, of a host is evaluated as a means of assessing the 35 "oxidative environment" of the host and the host's

susceptibility to VCAM-1 or redox-sensitive gene mediated disease.

In another embodiment, the level of circulating or cell-surface VCAM-1 or other appropriate marker 5 and the effect on that level of administration of an appropriate antioxidant is quantified.

In yet another assay, the sensitization of a host's vascular endothelial cells to polyunsaturated fatty acids or their oxidized 10 counterparts is evaluated. This can be accomplished, for example, by challenging a host with a PUFA or ox-PUFA and comparing the resulting concentration of cell-surface or circulating VCAM-1 or other surrogate marker to a population norm.

15 In another embodiment, in vivo models of atherosclerosis or other heart or inflammatory diseases that are mediated by VCAM-1 can be provided by administering to a host animal an excessive amount of PUFA or oxidized 20 polyunsaturated fatty acid to induce disease. These animals can be used in clinical research to further the understanding of these disorders.

In yet another embodiment of the invention, 25 compounds can be assessed for their ability to treat disorders mediated by VCAM-1 on the basis of their ability to inhibit the oxidation of a polyunsaturated fatty acid, or the interaction of a PUFA or ox-PUFA with a protein target.

This can be accomplished by challenging a host, 30 for example, a human or an animal such as a mouse, to a high level of PUFA or ox-PUFA and then determining the therapeutic efficacy of a test compound based on its ability to decrease circulating or cell surface VCAM-1 concentration. 35 Alternatively, an in vitro screen can be used that is based on the ability of the test compound to prevent the oxidation of a PUFA, or the interaction

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of a PUFA or ox-PUFA with a protein target in the presence of an oxidizing substance such as a metal, for example, copper, or an enzyme such as a peroxidase, lipoxygenase, cyclooxygenase, or 5 cytochrome P450.

In another embodiment, vascular endothelial cells are exposed to TNF- $\alpha$  or other VCAM-1 inducing material for an appropriate time and then broken by any appropriate means, for example by 10 sonication or freeze-thaw. The cytosolic and membrane compartments are isolated. Radiolabeled PUFA is added to defined amounts of the compartments. The ability of the liquid to convert PUFA to ox-PUFA in the presence or absence of a 15 test compound is assayed. Intact cells can be used in place of the broken cell system.

### III. Pharmaceutical Compositions

Humans, equine, canine, bovine and other animals, and in particular, mammals, suffering from 20 cardiovascular disorders, and other inflammatory conditions mediated by VCAM-1 or a redox sensitive gene can be treated by administering to the patient an effective amount of a compound that causes the removal, decrease in the concentration of, or 25 prevention of the formation of an oxidized polyunsaturated fatty acids, including but not limited to oxidized linoleic ( $C_{18} \Delta^{9,12}$ ), linolenic ( $C_{18} \Delta^{6,9,12}$ ), arachidonic ( $C_{20} \Delta^{5,8,11,14}$ ) and eicosatrienoic ( $C_{20} \Delta^{8,11,14}$ ) acids; other oxidation 30 signal; or other active compound, or a pharmaceutically acceptable derivative or salt thereof in a pharmaceutically acceptable carrier or diluent. The active materials can be administered by any appropriate route, for example, orally.

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parenterally, intravenously, intradermally, subcutaneously, or topically.

As used herein, the term pharmaceutically acceptable salts or complexes refers to salts or complexes that retain the desired biological activity of the above-identified compounds and exhibit minimal undesired toxicological effects. Nonlimiting examples of such salts are (a) acid addition salts formed with inorganic acids (for example, hydrochloric acid, hydrobromic acid, sulfuric acid, phosphoric acid, nitric acid, and the like), and salts formed with organic acids such as acetic acid, oxalic acid, tartaric acid, succinic acid, malic acid, ascorbic acid, benzoic acid, tannic acid, pamoic acid, alginic acid, polyglutamic acid, naphthalenesulfonic acid, naphthalenedisulfonic acid, and polygalacturonic acid; (b) base addition salts formed with polyvalent metal cations such as zinc, calcium, bismuth, barium, magnesium, aluminum, copper, cobalt, nickel, cadmium, sodium, potassium, and the like, or with an organic cation formed from N,N-dibenzylethylene-diamine, D-glucosamine, ammonium, tetraethylammonium, or ethylenediamine; or (c) combinations of (a) and (b); e.g., a zinc tannate salt or the like.

The active compound or a mixture of the compounds are administered in any appropriate manner, including but not limited to orally and intravenously. General range of dosage for any of the above-mentioned conditions will be from 0.5 to 500 mg/kg body weight with a dose schedule ranging from once every other day to several times a day. Preferred daily dosages are between approximately 1 and 3000 mg/patient/day, more preferably between approximately 5 and 500 mg/patient/day, and even

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more preferably, between approximately 25 and 500 mg/patient/day.

The active ingredient should be administered to achieve peak plasma concentrations of the active 5 compound of about 0.1 to 100  $\mu$ M, preferably about 1-10 $\mu$ M. This may be achieved, for example, by the intravenous injection of a solution or formulation of the active ingredient, optionally in saline, or an aqueous medium or administered as a bolus of the 10 active ingredient.

The compounds can also be administered directly to the vascular wall using perfusion balloon catheters following or in lieu of coronary or other arterial angioplasty. As an example, 2-5 mL of a 15 physiologically acceptable solution that contains approximately 1 to 500  $\mu$ M of the compound or mixture of compounds is administered at 1-5 atmospheres pressure. Thereafter, over the course of the next six months during the period of maximum 20 risk of restenosis, the active compounds are administered through other appropriate routes and dose schedules.

Relatively short term treatments with the active 25 compounds are used to cause the "shrinkage" of coronary artery disease lesions that cannot be treated either by angioplasty or surgery. A nonlimiting example of short term treatment is two to six months of a dosage ranging from 0.5 to 500 mg/kg body weight given at periods ranging from 30 once every other day to three times daily.

Longer term treatments can be employed to prevent the development of advanced lesions in high-risk patients. A long term treatment can extend for years with dosages ranging from 0.5 to 35 500 mg/kg body weight administered at intervals ranging from once every other day to three times daily.

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The active compounds can also be administered in the period immediately prior to and following coronary angioplasty as a means to reduce or eliminate the abnormal proliferative and 5 inflammatory response that currently leads to clinically significant re-stenosis.

The active compounds can be administered in conjunction with other medications used in the treatment of cardiovascular disease, including 10 lipid lowering agents such as probucol and nicotinic acid; platelet aggregation inhibitors such as aspirin; antithrombotic agents such as coumadin; calcium channel blockers such as varapamil, diltiazem, and nifedipine; angiotensin 15 converting enzyme (ACE) inhibitors such as captopril and enalopril, and  $\beta$ -blockers such as propanalol, terbutalol, and labetalol. The compounds can also be administered in combination with nonsteroidal antiinflammatories such as 20 ibuprofen, indomethacin, fenoprofen, mefenamic acid, flufenamic acid, sulindac. The compound can also be administered with corticosteroids.

The concentration of active compound in the drug composition will depend on absorption, 25 distribution, inactivation, and excretion rates of the drug as well as other factors known to those of skill in the art. It is to be noted that dosage values will also vary with the severity of the condition to be alleviated. It is to be further 30 understood that for any particular subject, specific dosage regimens should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the 35 compositions, and that the concentration ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the

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claimed composition. The active ingredient may be administered at once, or may be divided into a number of smaller doses to be administered at varying intervals of time.

5       Oral compositions will generally include an inert diluent or an edible carrier. They may be enclosed in gelatin capsules or compressed into tablets. For the purpose of oral therapeutic administration, the active compound can be 10 incorporated with excipients and used in the form of tablets, troches, or capsules. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition.

15       The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a 20 disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as 25 peppermint, methyl salicylate, or orange flavoring. When the dosage unit form is a capsule, it can contain, in addition to material of the above type, a liquid carrier such as a fatty oil. In addition, dosage unit forms can contain various 30 other materials which modify the physical form of the dosage unit, for example, coatings of sugar, shellac, or other enteric agents.

35       The active compound or pharmaceutically acceptable salt or derivative thereof can be administered as a component of an elixir, suspension, syrup, wafer, chewing gum or the like. A syrup may contain, in addition to the active

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compounds, sucrose as a sweetening agent and certain preservatives, dyes and colorings and flavors.

The active compound or pharmaceutically acceptable derivatives or salts thereof can also be administered with other active materials that do not impair the desired action, or with materials that supplement the desired action, such as antibiotics, antifungals, antiinflammatories, or 10 antiviral compounds.

Solutions or suspensions used for parenteral, intradermal, subcutaneous, or topical application can include the following components: a sterile diluent such as water for injection, saline 15 solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid; buffers such as acetates, citrates or phosphates and agents for the adjustment of tonicity such as sodium 20 chloride or dextrose. The parental preparation can be enclosed in ampoules, disposable syringes or 25 multiple dose vials made of glass or plastic.

Suitable vehicles or carriers for topical application are known, and include lotions, suspensions, ointments, creams, gels, tinctures, sprays, powders, pastes, slow-release transdermal 30 patches, aerosols for asthma, and suppositories for application to rectal, vaginal, nasal or oral mucosa.

Thickening agents, emollients, and stabilizers can be used to prepare topical compositions. 35 Examples of thickening agents include petrolatum, beeswax, xanthan gum, or polyethylene glycol, humectants such as sorbitol, emollients such as

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mineral oil, lanolin and its derivatives, or squalene. A number of solutions and ointments are commercially available.

5 Natural or artificial flavorings or sweeteners can be added to enhance the taste of topical preparations applied for local effect to mucosal surfaces. Inert dyes or colors can be added, particularly in the case of preparations designed for application to oral mucosal surfaces.

10 The active compounds can be prepared with carriers that protect the compound against rapid release, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can 15 be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Many methods for the preparation of such formulations are patented or generally known to those skilled in the 20 art.

If administered intravenously, preferred carriers are physiological saline or phosphate buffered saline (PBS).

25 The active compound can also be administered through a transdermal patch. Methods for preparing transdermal patches are known to those skilled in the art. For example, see Brown, L., and Langer, R., *Transdermal Delivery of Drugs*, Annual Review of Medicine, 39:221-229 (1988), incorporated herein by 30 reference.

In another embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including 35 implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides,

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5 polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc.

10 Liposomal suspensions may also be pharmaceutically acceptable carriers. These may be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811 (which is incorporated herein by reference in its entirety). For example, liposome formulations may be prepared by dissolving appropriate lipid(s) (such as stearoyl phosphatidyl 15 ethanolamine, stearoyl phosphatidyl choline, arachadoyl phosphatidyl choline, and cholesterol) in an inorganic solvent that is then evaporated, leaving behind a thin film of dried lipid on the surface of the container. An aqueous solution of 20 the active compound or its monophosphate, diphosphate, and/or triphosphate derivatives are then introduced into the container. The container is then swirled by hand to free lipid material from the sides of the container and to disperse lipid 25 aggregates, thereby forming the liposomal suspension.

30 Modifications and variations of the present invention will be obvious to those skilled in the art from the foregoing detailed description of the invention. Such modifications and variations are intended to come within the scope of the appended

We claim.

1. A method for supressing the expression of VCAM-1 comprising administering an effective amount of a substance that prevents or minimizes the oxidation of a polyunsaturated fatty acid.

2. A method for suppressing the expression of a redox-sensitive gene comprising administering an effective amount of a substance that prevents or minimizes the oxidation of a polyunsaturated fatty acid.

3. A method for activating a gene that is suppressed by the oxidation of a polyunsaturated fatty acid, comprising administering an effective amount of a substance that prevents or minimizes the oxidation of a polyunsaturated fatty acid.

4. A method for suppressing the expression of VCAM-1 comprising administering an effective amount of a substance that prevents the interaction between a polyunsaturated fatty acid and a protein that mediates the expression of VCAM-1.

5. The method of claims 1-4, wherein the polyunsaturated acid is selected from the group consisting of oxidized linoleic ( $C_{18} \Delta^{9,12}$ ), linolenic ( $C_{18} \Delta^{6,9,12}$ ), arachidonic ( $C_{20} \Delta^{5,8,11,14}$ ) and eicosatrienoic ( $C_{20} \Delta^{8,11,14}$ ) acid.

6. The method of claim 2 or 3, wherein the redox-sensitive gene is selected from the group consisting of those expressing cytokines involved in initiating the immune response (e.g., IL-1 $\beta$ ), chemoattractants that promote the migration of inflammatory cells to a point of injury (e.g., MCP-1), growth factors (e.g., IL-6 and the thrombin receptor), and adhesion molecules (e.g., VCAM-1 and E-selectin).

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7. The method of claims 1-4, wherein the substance is pyrrolidine dithiocarbamate, or its pharmaceutically acceptable salt.

8. A method for the prediction or assessment of disorders mediated by VCAM-1 in vivo, comprising quantifying the level of oxidized polyunsaturated fatty acid in the tissue or blood.

9. A method for the prediction or assessment of redox-sensitive gene mediated disease in vivo, comprising quantifying the level of oxidized polyunsaturated fatty acid in the tissue or blood.

11. A method for the prediction or assessment of disorders mediated by VCAM-1 in vivo, comprising quantifying a surrogate marker for the level of oxidized polyunsaturated fatty acid in the tissue or blood.

12. A method for the prediction or assessment of redox-sensitive gene mediated disease in vivo, comprising quantifying a surrogate marker for the level of oxidized polyunsaturated fatty acid in the tissue or blood.

13. The method of claim 11, wherein the surrogate marker is circulating or cell-surface VCAM-1.

14. A method for the evaluation of the sensitization of a host's vascular endothelial cells to polyunsaturated fatty acids or their oxidized counterparts, comprising challenging a host with a PUFA or ox-PUFA and comparing the resulting concentration of cell-surface or circulating VCAM-1 or other surrogate marker to a population norm.

15. A method to screen compounds for their ability to treat disorders mediated by VCAM-1 comprising evaluating the ability of the compound to inhibit the oxidation of a polyunsaturated fatty acid.

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16. A method to screen compounds for their ability to treat disorders mediated by VCAM-1 comprising evaluating the ability of the compound to inhibit the interaction of a PUFA or ox-PUFA with a protein target.

17. A method for the treatment of a cardiovascular disease in humans comprising administering an effective amount of a dithiocarbamate of the formula A-SC(S)-B;

wherein A selected from the group consisting of hydrogen, a pharmaceutically acceptable cation, and a physiologically cleavable leaving group;

and B is selected from the group consisting of alkyl, alkenyl, alkynyl, alkaryl, aralkyl, haloalkyl, haloalkenyl, haloalkynyl, aryl, alkaryl, hydrogen, C<sub>1-6</sub> alkoxy-C<sub>1-10</sub> alkyl, C<sub>1-6</sub> alkylthio-C<sub>1-10</sub> alkyl, NR<sup>2</sup>R<sup>3</sup>, -(CHOH)<sub>n</sub>CH<sub>2</sub>OH, wherein n is 0, 1, 2, 3, 4, 5, or 6, -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>1</sup>, including alkylacetyl, alkylpropionyl, and alkylbutyryl, and hydroxy(C<sub>1-6</sub>)alkyl-.

18. The method of claim 17, wherein A is hydrogen or a pharmaceutically acceptable cation selected from the group consisting of sodium, potassium, calcium, magnesium, aluminum, zinc, bismuth, barium, copper, cobalt, nickel, or cadmium.

19. The method of claim 17, wherein A is a salt-forming organic acid.

20. The method of claim 19, wherein A is selected from the group consisting of choline, acetic acid, oxalic acid, tartaric acid, succinic acid, malic acid, ascorbic acid, benzoic acid, tannic acid, pamoic acid, alginic acid, polyglutamic acid, naphthalenesulfonic acid, naphthalenedisulfonic acid, and polygalacturonic acid.

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21. The method of claim 17, wherein A is a cation formed from ammonia or other nitrogenous base.

22. The method of claim 21, wherein A is a nitrogenous heterocycle, or a moiety of the formula  $NR^4R^5R^6R^7$ , wherein  $R^4$ ,  $R^5$ ,  $R^6$ , and  $R^7$  are independently hydrogen,  $C_{1-6}$  alkyl, hydroxy( $C_{1-6}$ )alkyl, aryl, N,N-dibenzylethylene-diamine, D-glucosamine, tetraethylammonium, or ethylenediamine.

23. The method of claim 17, wherein A is a physiologically cleavable leaving group.

24. The method of claim 17, wherein A is an acyl group.

25. The method of claim 17, wherein B is  $NR^2R^3$ , wherein  $R^2$  and  $R^3$  are selected from the group consisting of alkyl;  $-(CHOH)_n(CH_2)_nOH$ , wherein n is 0, 1, 2, 3, 4, 5, or 6;  $-(CH_2)_nCO_2R^1$ ,  $-(CH_2)_nCO_2R^4$ ; hydroxy( $C_{1-6}$ )alkyl-; alkenyl; alkyl( $CO_2H$ ), alkenyl( $CO_2H$ ), alkynyl( $CO_2H$ ), or aryl, or  $R^2$  and  $R^3$  can together constitute a bridge of the formula  $-(CH_2)_m-$ , wherein m is 3, 4, 5, or 6, and wherein  $R^4$  is selected from the group consisting of aryl, alkaryl, or aralkyl, including acetyl, propionyl, and butyryl.

26. The method of claim 17, wherein B is a heterocyclic or alkylheterocyclic group.

27. The method of claim 26, wherein the heterocycle is partially or totally hydrogenated.

28. The method of claim 17, wherein B is the residue of a pharmaceutically-active compound or drug which is directly linked to A-SC(S)- or linked through a divalent linking moiety.

29. The method of claim 17, wherein B is selected from the group consisting of probucol, nicotinic acid, aspirin, coumadin, varapamil, diltiazem, nifedipine, captopril, enalopril, propanalol, terbutalol, labetalol, ibuprofen,

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indomethacin, fenoprofen, mefenamic acid, flufenamic acid, sulindac, and a corticosteroid.

30. The method of claim 17, wherein the dithiocarbamate is an amino acid derivative of the structure  $AO_2C-R^9-NR^{10}-C(S)SA$ , wherein  $R_9$  is B or the internal residue of an amino acid and  $R^{10}$  is hydrogen or lower alkyl.

31. The method of claim 17, wherein B is a polymer to which one or more dithiocarbamate groups are attached, either directly, or through any suitable linking moiety.

32. The method of claim 17, wherein the polymer is biodegradable.

33. The method of claim 32, wherein the polymer is selected from the group consisting of peptides, proteins, nucleoproteins, lipoproteins, glycoproteins, synthetic and natural polypeptides and polyamino acids, polyorthoesters, poly( $\alpha$ -hydroxy acids), polyanhydrides, polysaccharides, and polycaprolactone.

34. The method of claim 1, wherein B-C(S)S- is pyrrolidine-N-carbodithioate.

35. The method of claim 17 wherein the cardiovascular disease is atherosclerosis.

36. The method of claim 17, wherein the cardiovascular disease is post-angioplasty restenosis.

37. The method of claim 17, wherein the cardiovascular disease is coronary artery disease.

38. The method of claim 17, wherein the cardiovascular disease is angina.

39. The method of claim 17, wherein the cardiovascular disease is a small vessel disease.

40. The method of claim 17, wherein the dithiocarbamate is administered in a dosage of between 0.5 and 500 mg/kg body weight.

41. The method of claim 17, wherein the dithiocarbamate is administered by perfusion balloon catheter.

42. The method of claim 17, wherein the dithiocarbamate is administered in combination with a pharmaceutical agent selected from the group consisting of a lipid lowering agent, a platelet aggregation inhibitor, an antithrombotic agent, a calcium channel blocker, an angiotensin converting enzyme (ACE) inhibitor, a  $\beta$ -blocker, a nonsteroidal antiinflammatory, and a corticosteroid.

43. A method for the suppression of VCAM-1 expression in human cells comprising administering an effective amount of the dithiocarbamate described in claim 17.

44. A method for the treatment of an inflammatory skin disease that is mediated by VCAM-1 comprising administering an effective amount of the dithiocarbamate described in claim 17.

45. A method for the treatment of a human endothelial disorder that is mediated by VCAM-1 comprising administering an effective amount of the dithiocarbamate described in claim 17.

46. The method of claim 45, wherein the disorder is selected from the group consisting of asthma, psoriasis, eczematous dermatitis, Kaposi's sarcoma, multiple sclerosis, and proliferative disorders of smooth muscle cells.

47. A method for the treatment of an inflammatory condition that is mediated by a mononuclear leucocyte comprising administering an effective amount of the dithiocarbamate described in claim 17.

48. The dithiocarbamate disclosed in any of claims 17-34.

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49. A pharmaceutical composition comprising an effective amount to treat cardiovascular disease of a compound disclosed in any of claims 17-34.

50. A pharmaceutical composition comprising an effective amount to treat a disorder mediated by VCAM-1 of a compound disclosed in any of claims 17-34.

**Effects of Linoleic Acid and 13-HPODE on the Kinetics of  
VCAM-1 Gene Expression**

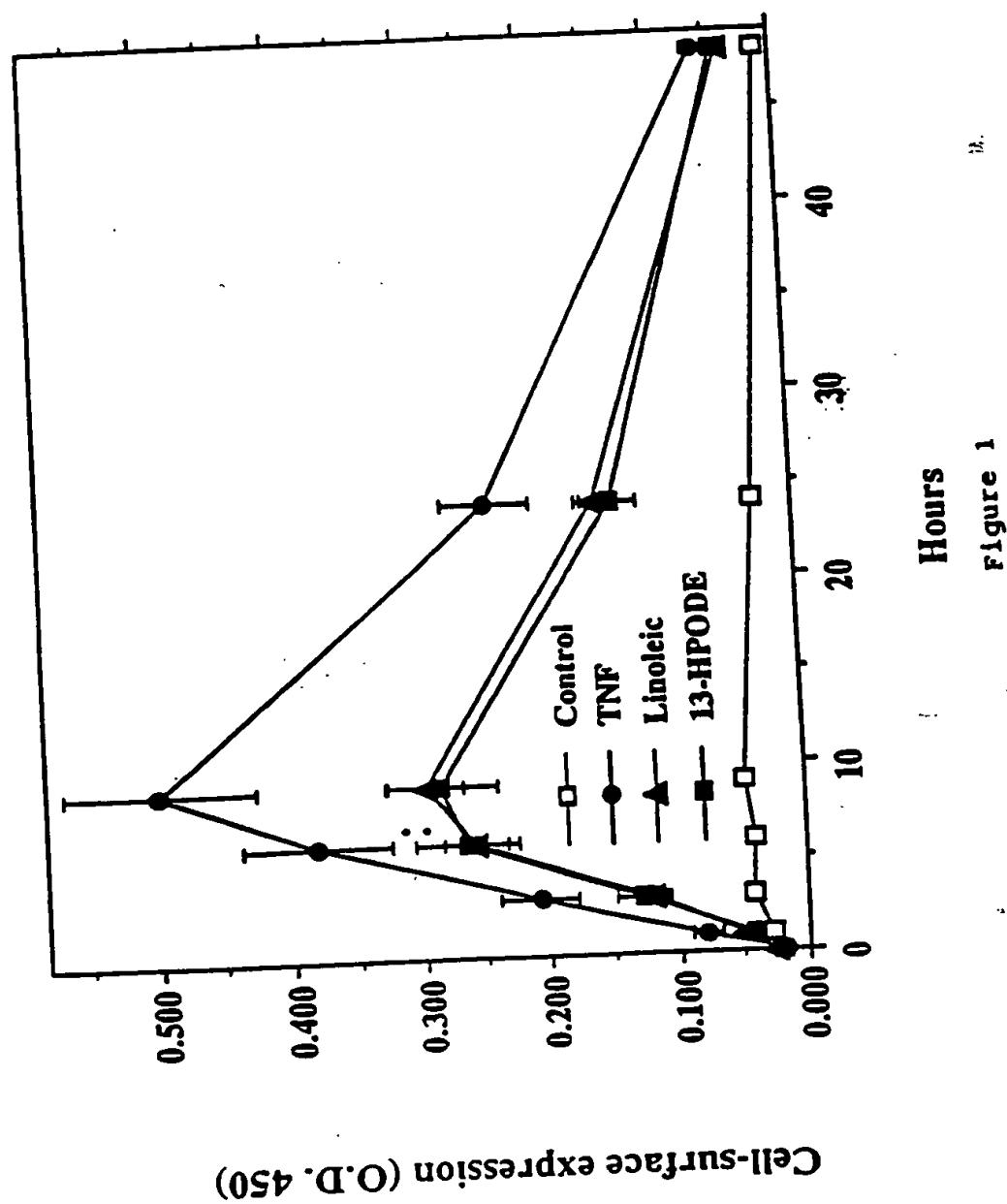


Figure 1



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## Effect of Linoleic Acid and 13-HPODE Dose Response on VCAM-1 Gene Expression

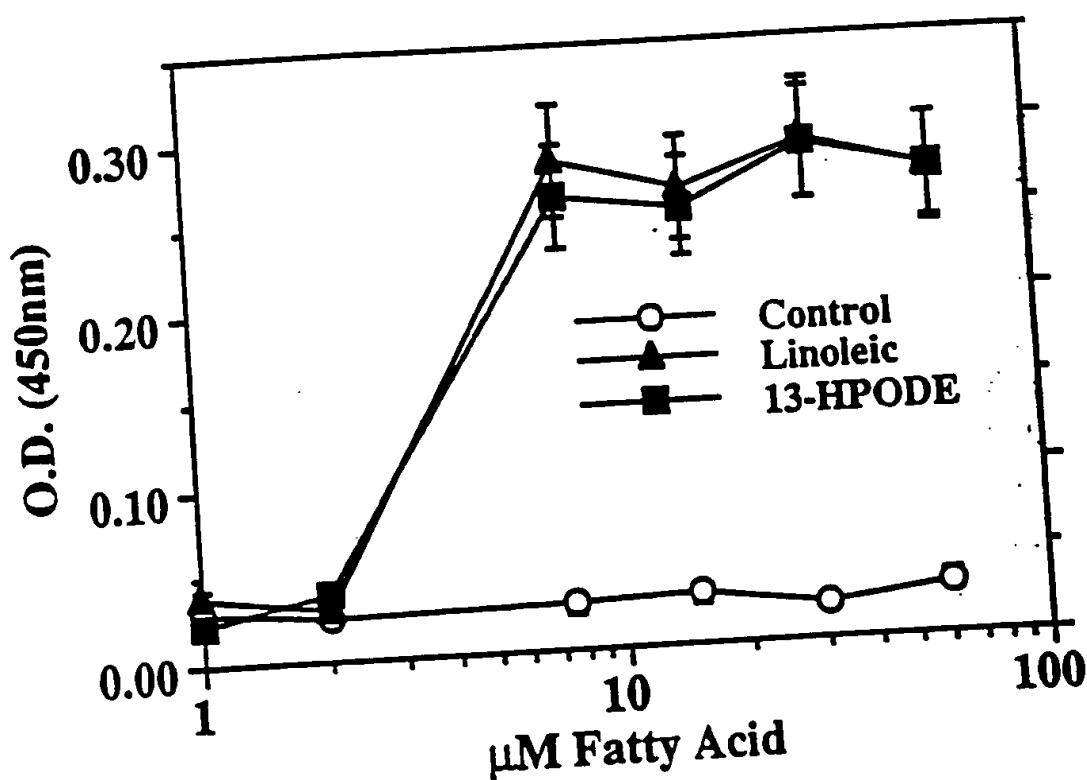


Figure 2



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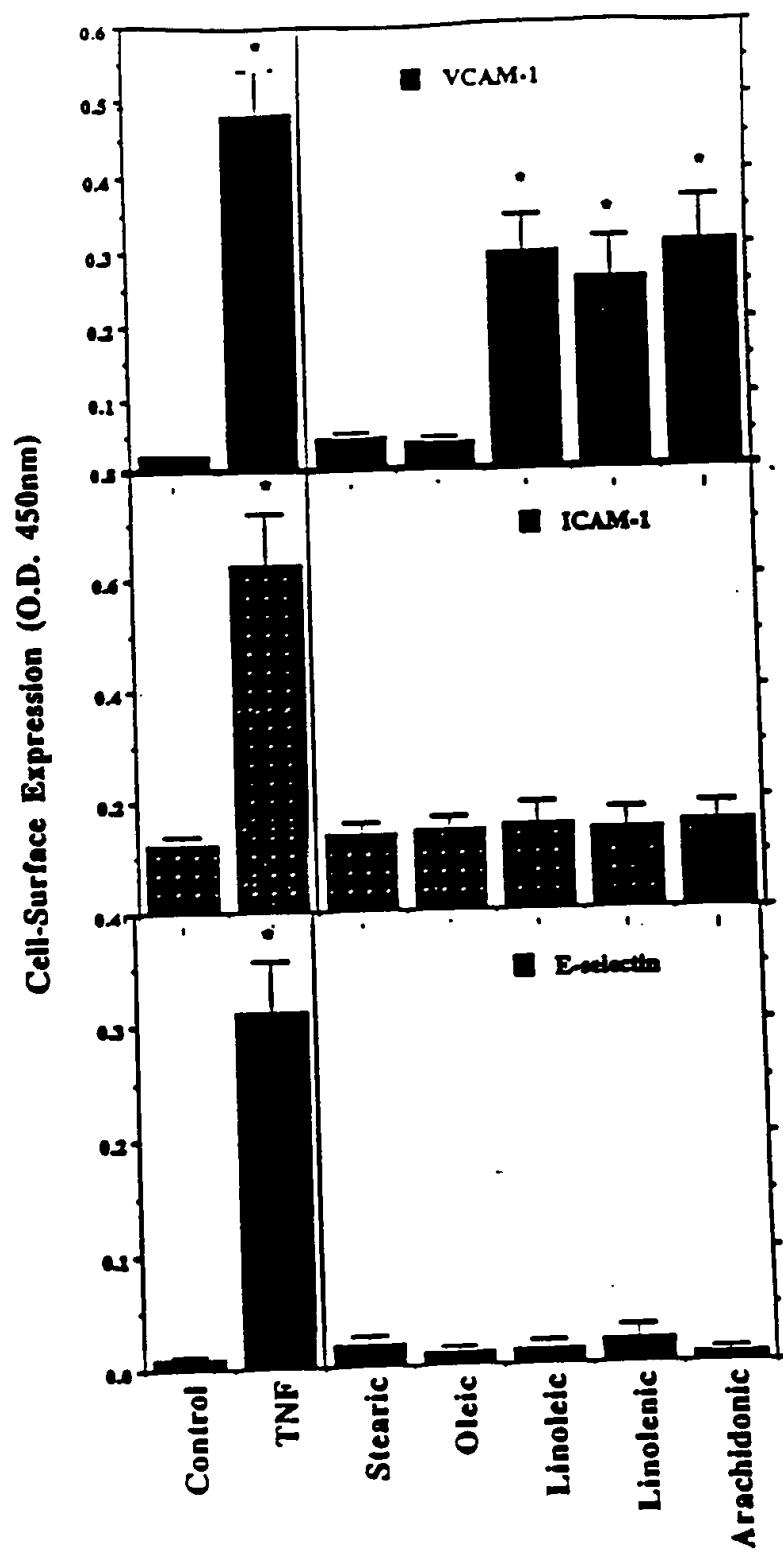
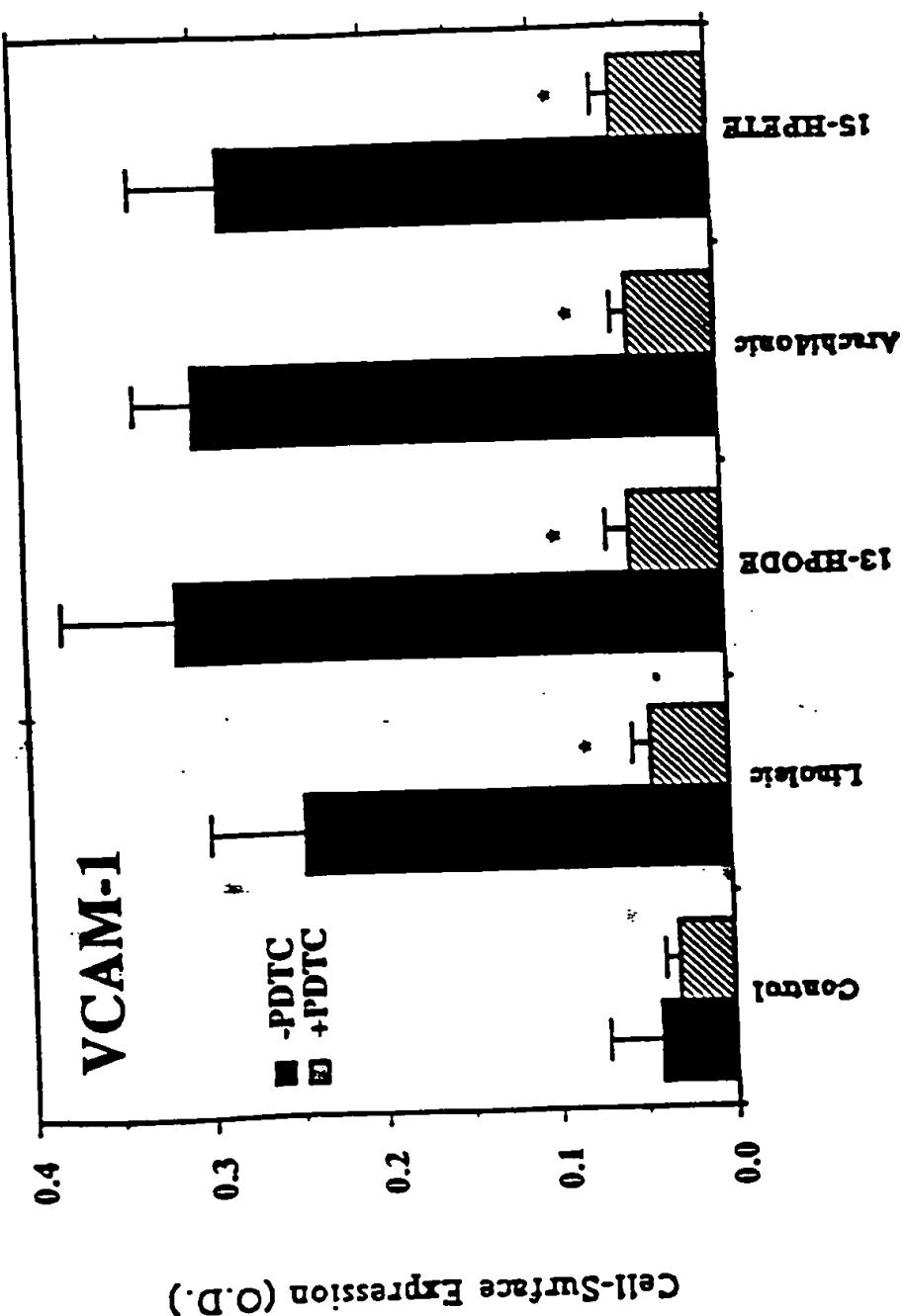


Figure 3



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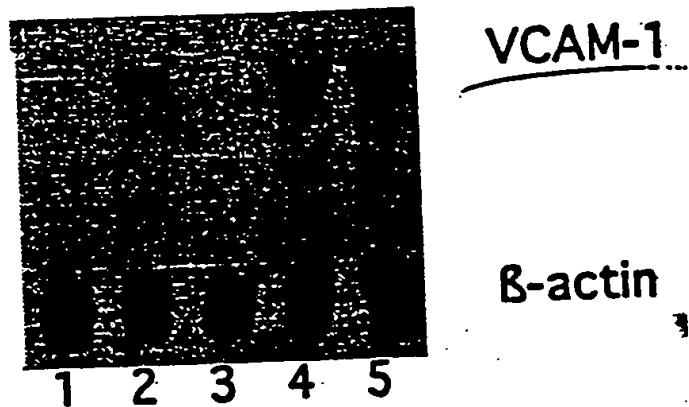


Figure 5



6 / 2 0

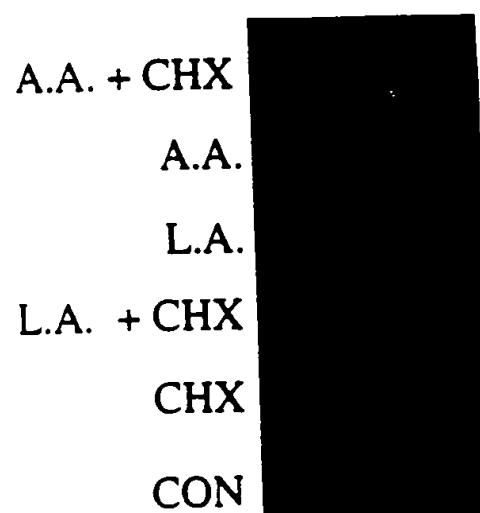
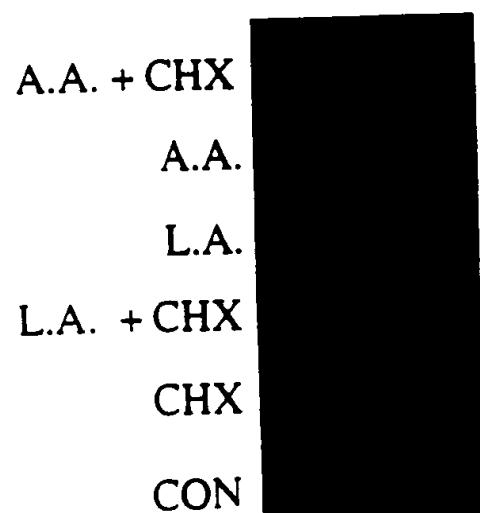
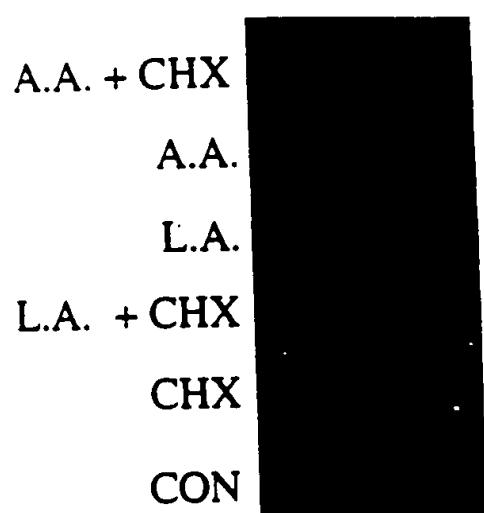
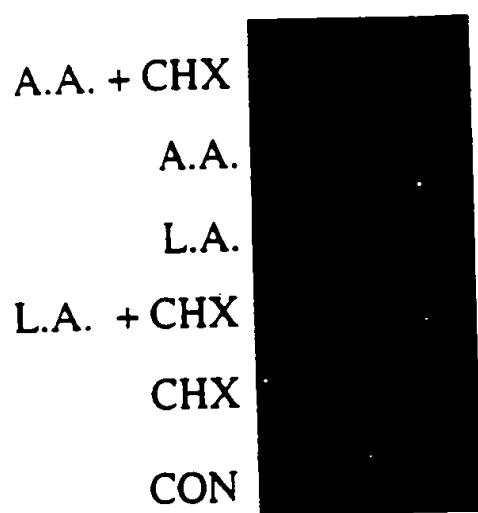


Figure 6



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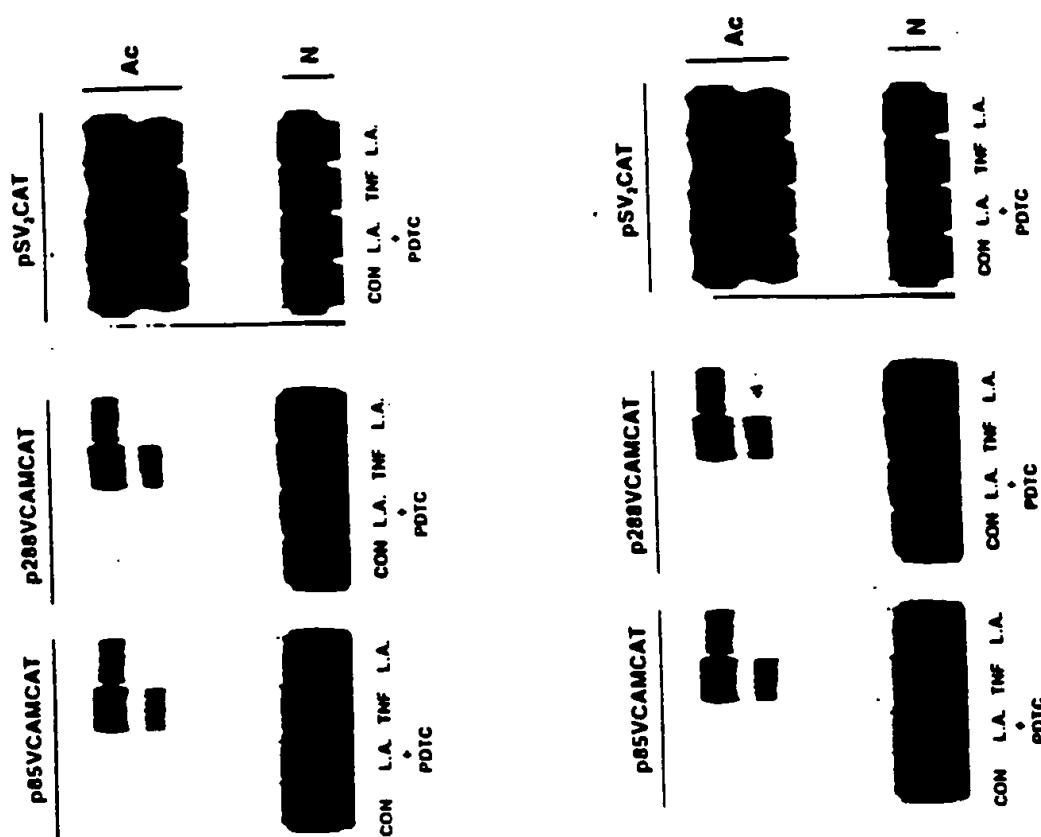


Figure 7



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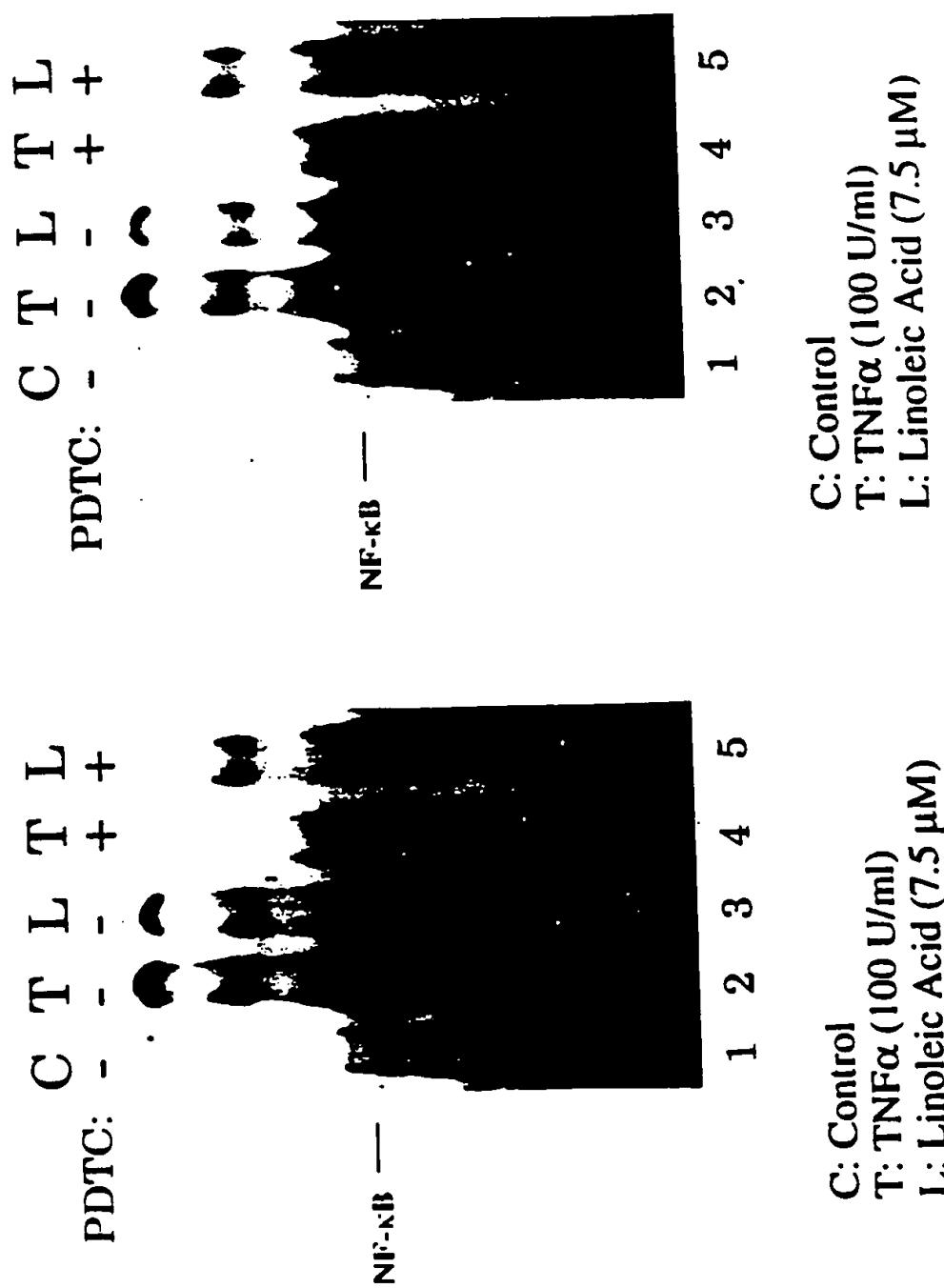
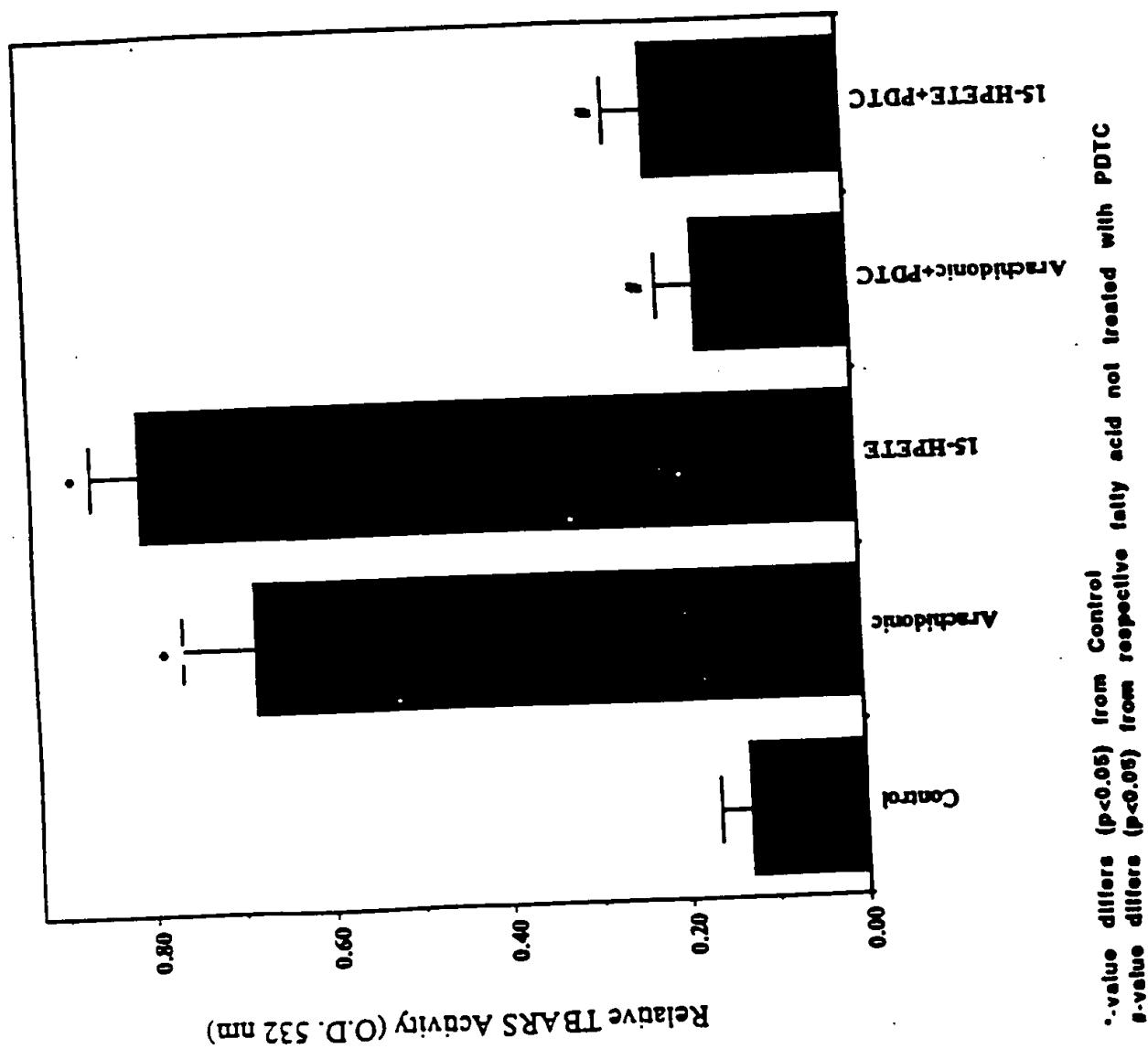


Figure 8

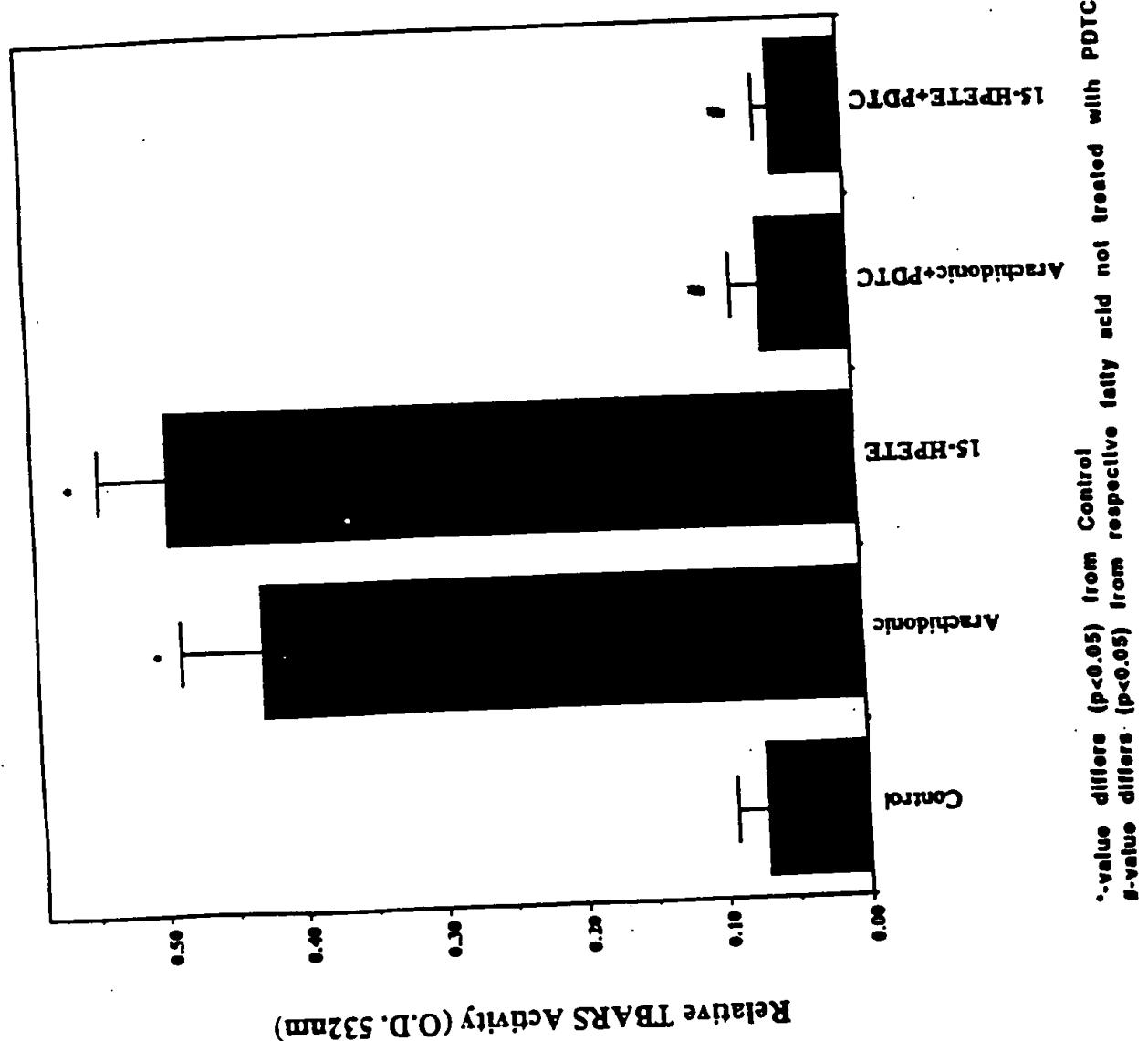


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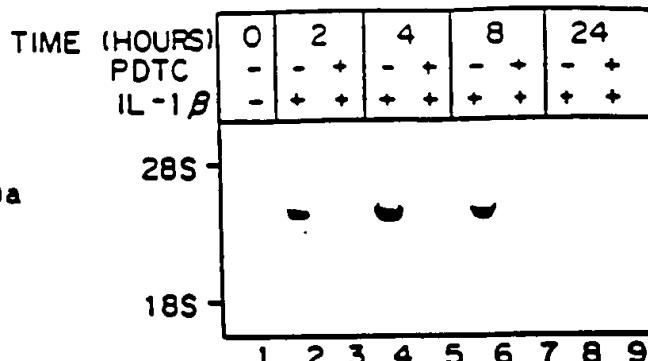


Figure 10a

VCAM-1

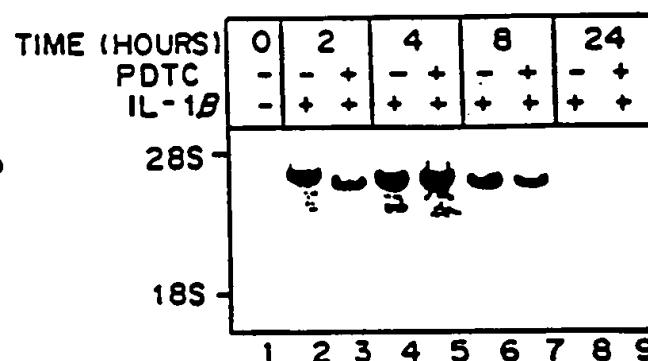


Figure 10b

E-SELECTIN

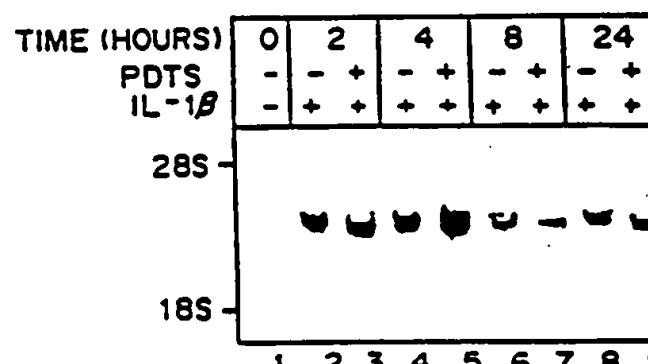
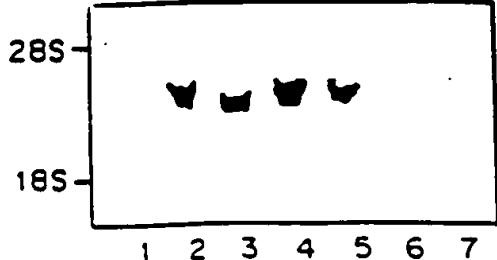


Figure 10c

ICAM-1

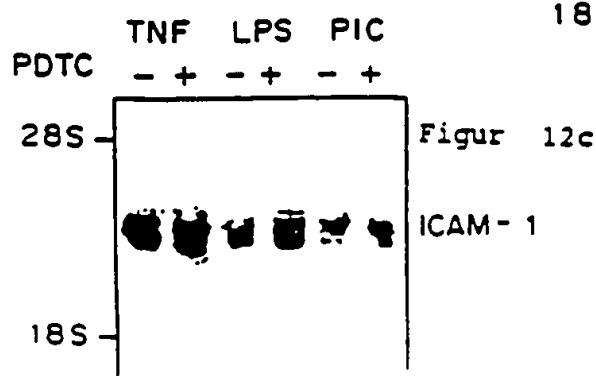
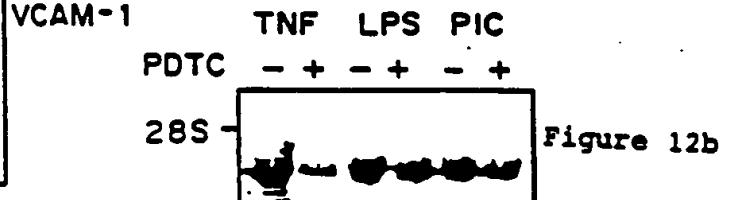
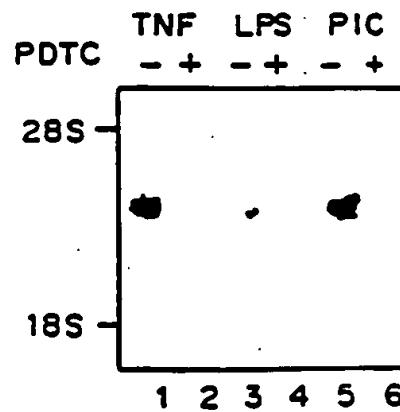
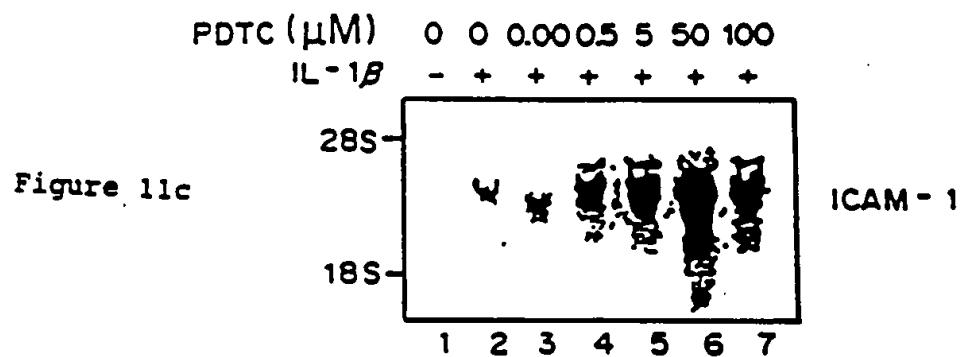
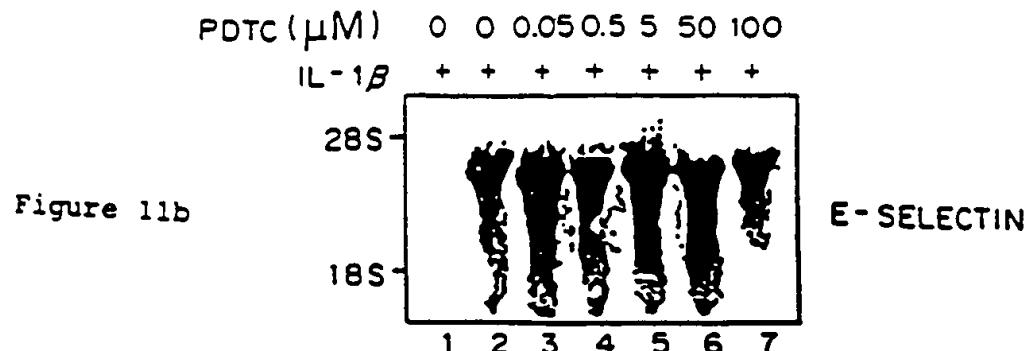
Figure 11a

PDT<sub>C</sub> ( $\mu$ M) 0 0 0.05 0.5 5 50 100IL-1 $\beta$  - + + + + +

VCAM-1



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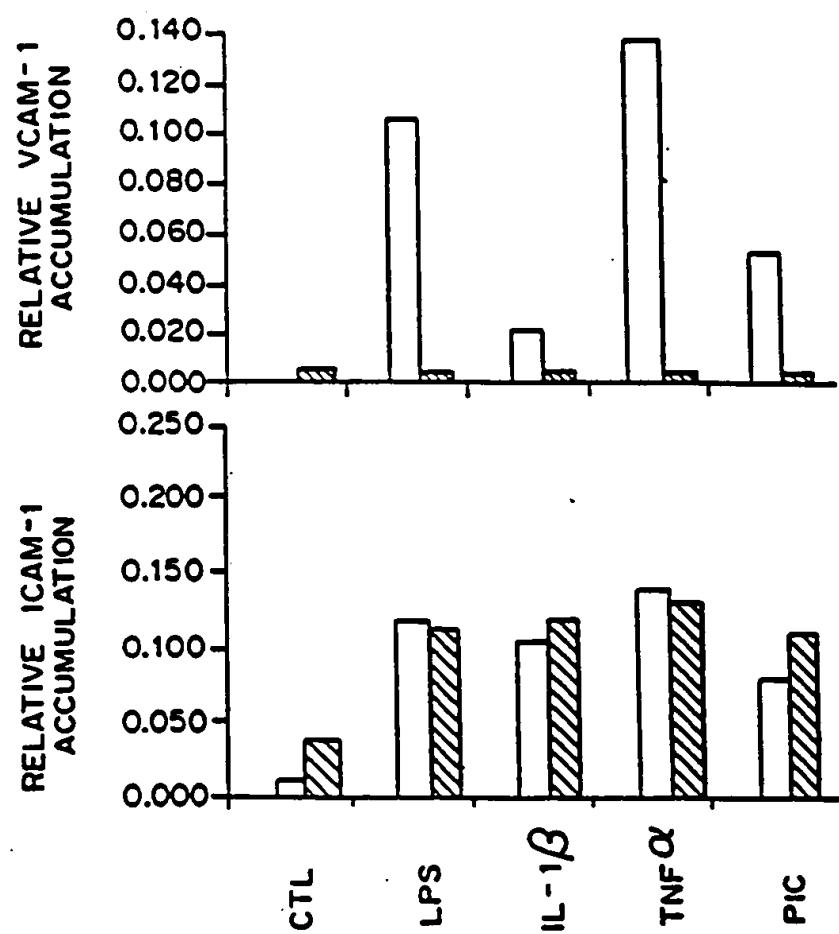


Figure 13



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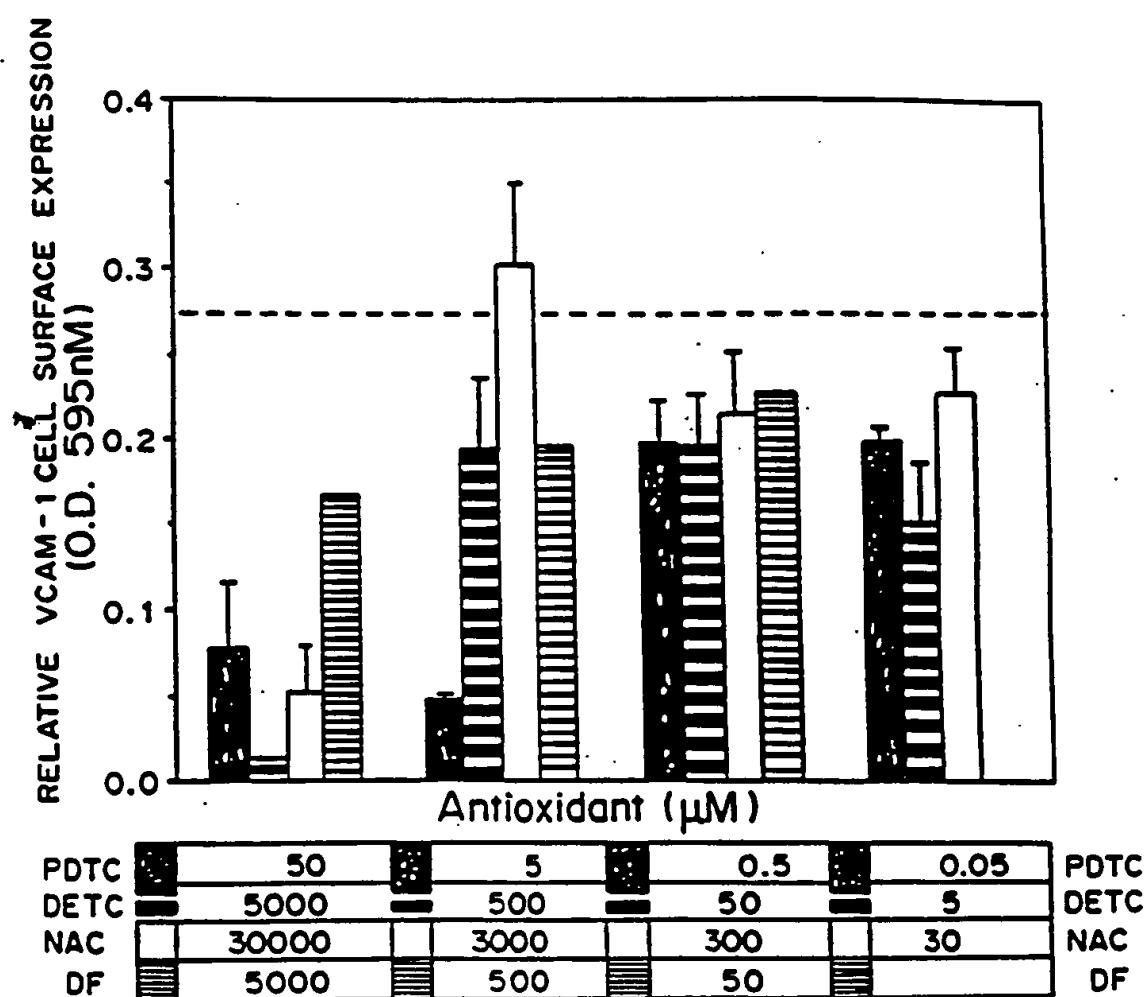


Figure 14



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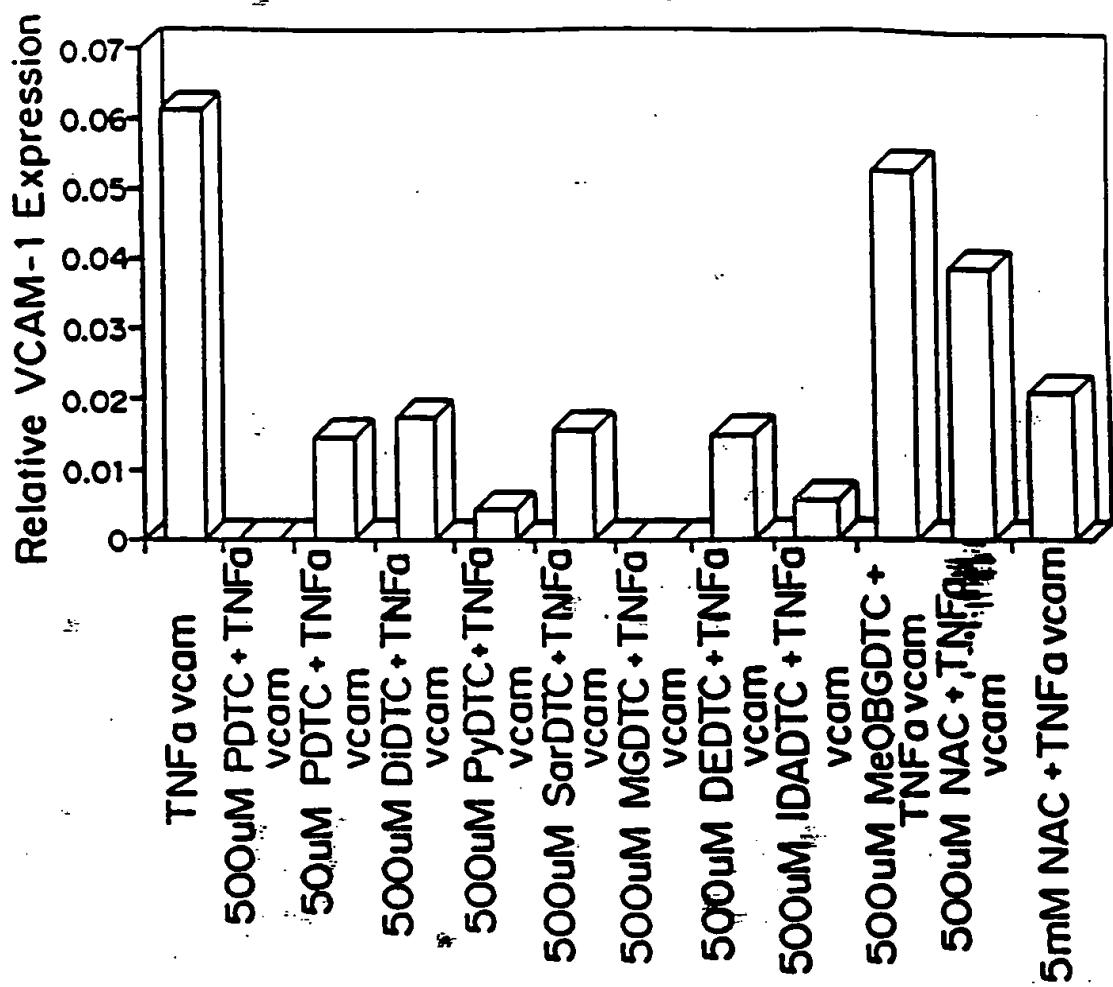
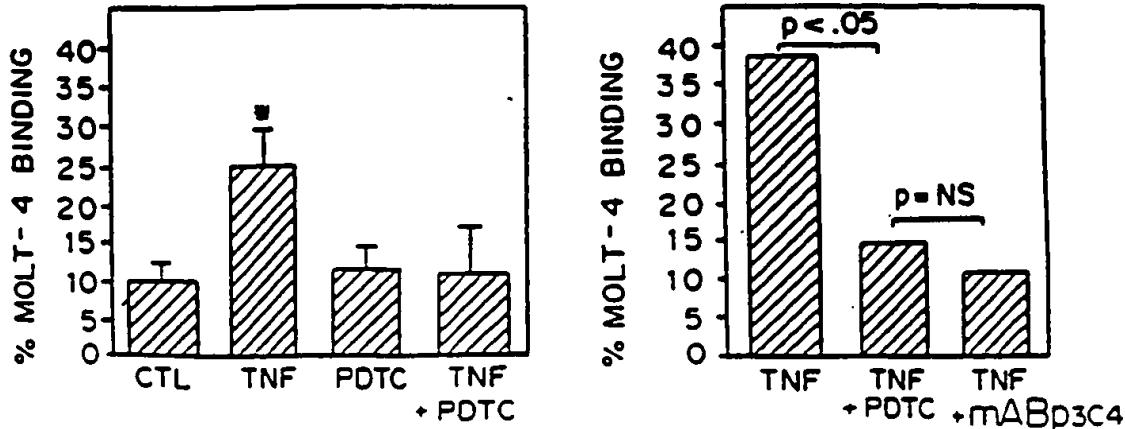


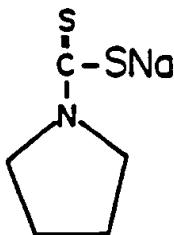
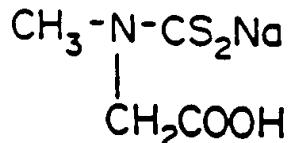
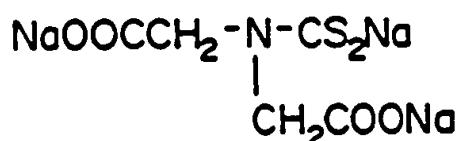
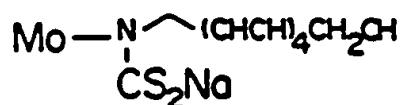
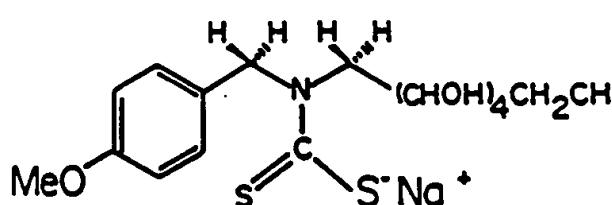
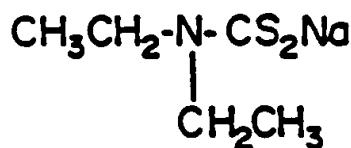
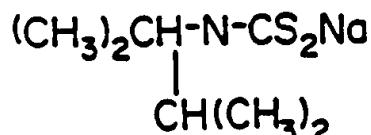
Figure 15





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Figure 17

SODIUM PYRROLIDINE-*N*-CARBODITHIOATESODIUM *N*-METHYL-*N*-CARBOXYMETHYL-*N*-CARBODITHIOATE  
(OR SODIUM SARCOSINEDITHIOCARBAMATE)TRISODIUM *N,N*-di(CARBOXYMETHYL)-*N*-CARBODITHIOATE  
(OR IMINODIACETIC ACID DITHIOCARBAMATE,  
TRISODIUM)SODIUM *N*-METHYL-*D*-GLUCAMINE-*N*-CARBODITHIOATESODIUM *N*-(4-METHOXYBENZYL)-*D*-GLUCAMINE-*N*-CARBODITHIOATESODIUM *N,N*-DIETHYL-*N*-CARBODITHIOATE (OR SODIUM DIETHYLDITHIOCARBAMATE)SODIUM *N,N*-DIISOPROPYL-*N*-CARBODITHIOATE  
(OR SODIUM DIISOPROPYLDITHIOCARBAMATE)



### Effect of PDTc on the formation of fluorescent adducts

(1 Micromol of 13-HPODE was incubated with 200 micrograms of BSA in the presence of PDTc for 6 days. Fluorescence was measured at 430-460 nm with excitation set at 330-360 nm)

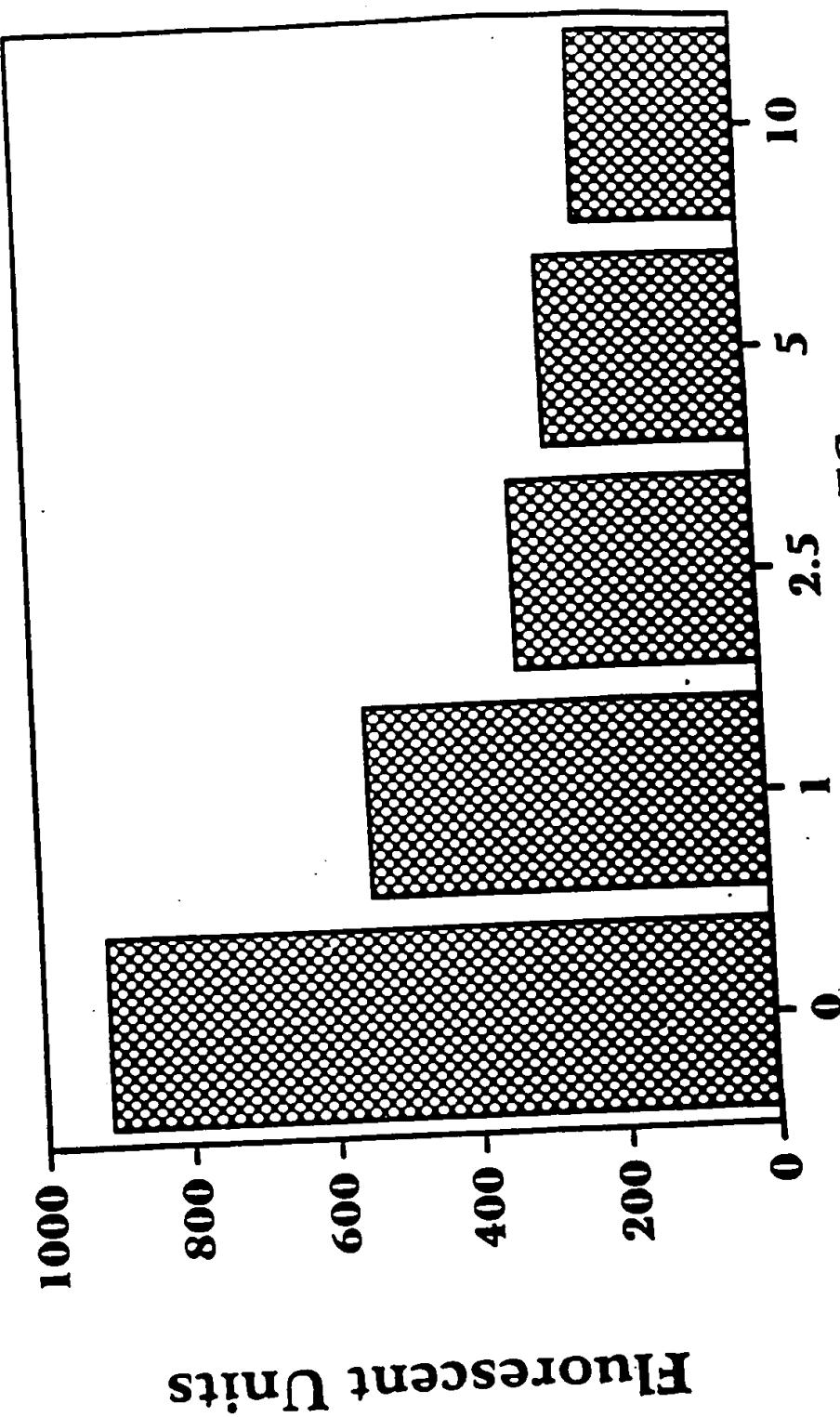


Figure 18



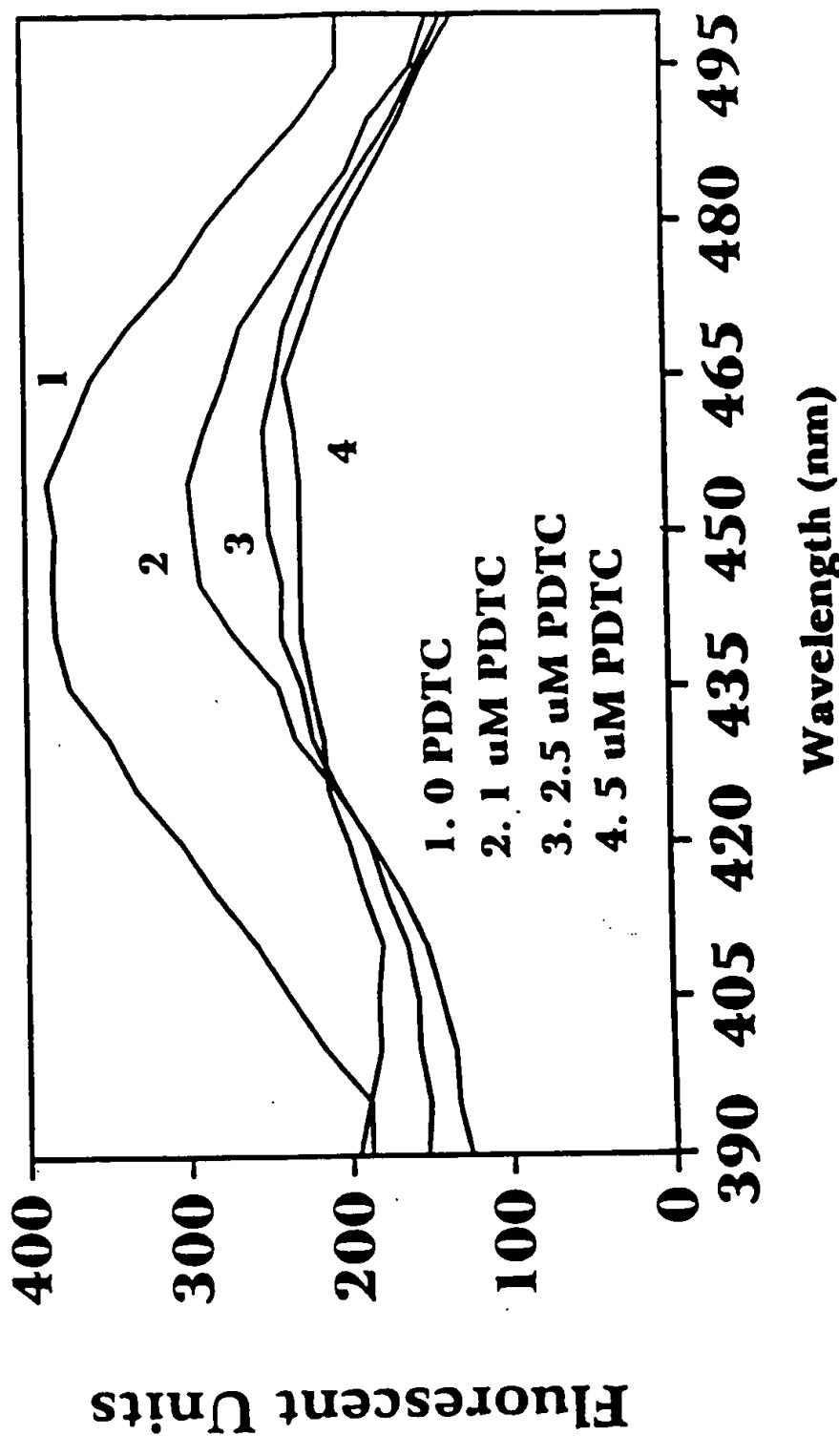
**Effect of PDTc on the formation of fluorescent adducts from BSA and LOOH**

Figure 19



### Effect of PDTC on the Oxidation of LDL by HRP

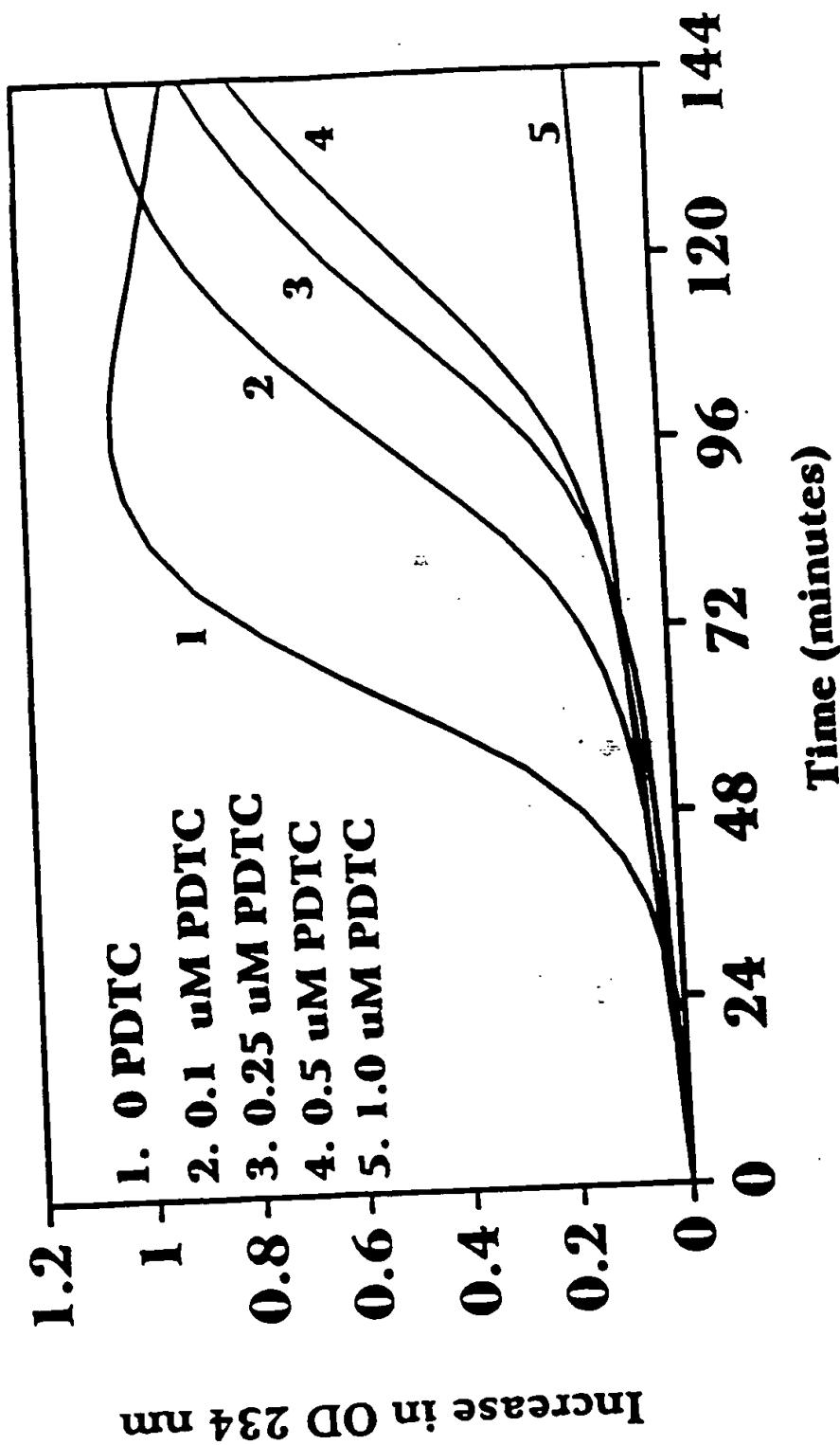


Figure 20



Cells	Linoleic	Ox-linoleic
Control	9561	11445
TNF $\alpha$	8874	27894
IL-1 $\beta$	9134	25813*
TNF $\alpha$ +PDTC	8444	7966#
IL-1 $\beta$ +PDTC	8257	8012#

\*-value differs ( $p<0.05$ ) from Control

#-value differs ( $p<0.05$ ) from corresponding group not treated with PDTC



## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US95/05880

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :Please See Extra Sheet.

US CL :Please See Extra Sheet.

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : Please See Extra Sheet.

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS ONLINE, BIOSIS, MEDLINE, APS

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	LABORATORY INVESTIGATION, Volume 64, No. 3, issued 1991, Pober et al., "What Can Be Learned From the Expression of Endothelial Adhesion Molecules in Tissues?," pages 301-305, see entire document.	1-6, 11-13, 43-50
X	INTERNATIONAL JOURNAL OF IMMUNOPHARMACOLOGY, Volume 6, No. 3, issued 1984, Corke et al., "The Influence of Diethyl-dithiocarbamate ('Imuthiol') on Mononuclear Cells in Vitro," pages 245-247, see entire document.	48-50
X	US, A, 4,870,101 (KU ET AL.) 26 September 1989, see entire document.	48-50
Y		1-6, 11-13, 43-50

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	
*A*	document defining the general state of the art which is not considered to be part of particular relevance
*E*	earlier document published on or after the international filing date
*L*	document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
*O*	document referring to an oral disclosure, use, exhibition or other means
*P*	document published prior to the international filing date but later than the priority date claimed
"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"Z"	document member of the same patent family

Date of the actual completion of the international search

04 SEPTEMBER 1995

Date of mailing of the international search report

13 SEP 1995

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*Peter G. O'Sullivan*  
PETER G. O'SULLIVAN  
Telephone No. (703) 308-1235

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US95/05880

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Chemical Abstracts, 119:23788, issued 1993, Pasqualini et al., "Brain-tropic Radiopharmaceutical Compounds Comprising a Transition Metal Nitride Complex, and Preparation Method Therefor," see abstract No. 23788, WO 9301839.	48-50
Y	US, A, 4,670,471 (CLARK) 02 June 1987, see abstract.	1-6, 11-13, 43-50
A	THE NEW ENGLAND JOURNAL OF MEDICINE, Volume 320, Number 14, issued 1989, Steinberg et al., "Beyond Cholesterol: Modifications of Low-Density Lipoprotein That Increase Its Atherogenicity, pages 915-924.	1-50
X	Chemical Abstracts, Volume 81, issued 1974, Tiwari et al., "Possible Anti-Parkinsonian Compounds IV. Synthesis of Piperazine bis[amino(thiocarbonyl)thio-acetates and propionates]," see page 527, column 1, abstract No. 105447b.	48-50
Y	THE JOURNAL OF CLINICAL INVESTIGATION, Volume 77, No. 2, issued 1986, Parthasarathy et al., "Probucol Inhibits Exudative Modification of Low Density Lipoprotein," pages 641-644.	48-50
X	THE LANCET, Volume II, No. 8613, issued 1988, Lang et al., "Randomized, Double-Blind, Placebo-Controlled Trial of Dithiocarb Sodium ('Imuthiol') In Human Immunodeficiency Virus Infection," pages 702-706, see entire document.	48-50
Y		48-50

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US95/05880

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
  
2.  Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.  
 No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US95/05880

**A. CLASSIFICATION OF SUBJECT MATTER:**  
IPC (6):

A61K 31/21, 31/40, 38/06; C07C 327/00, 327/36, 333/16, 333/20, 333/24, 333/32; C07D 207/04; C07K 5/09

**A. CLASSIFICATION OF SUBJECT MATTER:**  
US CL :

514/18, 423, 478, 479, 484, 485, 487, 488, 489, 506, 513, 824, 825, 826, 861, 863, 886; 530/331; 548/431; 558/230, 235; 564/76; 568/21, 25

**B. FIELDS SEARCHED**

Minimum documentation searched

Classification System: U.S.

514/18, 423, 478, 479, 484, 485, 487, 488, 489, 506, 513, 824, 825, 826, 861, 863, 886; 530/331; 548/431; 558/230, 235; 564/76; 568/21, 25

**BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING**

This ISA found multiple inventions as follows:

- I. Claims 1-50, drawn to heterocyclic dithiocarbamate compounds, compositions and methods.
- II. Claims 1-6, 8-25, 28-30, 35-50, drawn to non-polymeric, non-heterocyclic dithiocarbamate compounds, compositions and methods.
- III. Claims 1-6, 8-25, 28-30, 35-50, drawn to non-polymeric, non-heterocyclic thiuram disulfide compounds, compositions and methods.
- IV. Claims 1-6, 8-24, 31-33, 35-50, drawn to polymeric dithiocarbamate compounds, compositions and methods.
- V. Claims 1-6, 9-17, 23, 30, 35-50, drawn to dithiocarboxylate ester compounds, compositions and methods.
- VI. Claims 1-6, 8-17, 23, 25, 28, 35-50, drawn to bis thiocarbonylsulfide compounds, compositions and methods.

The inventions listed as Groups I-VI do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: the active compounds of each group are structurally diverse from the other groups so that references for one group would not necessarily suggest the others to one skilled in the art.